



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 :  C12N		A2	(11) International Publication Number: <b>WO 00/39279</b>  (43) International Publication Date: 6 July 2000 (06.07.00)
<p>(21) International Application Number: PCT/US99/31169</p> <p>(22) International Filing Date: 30 December 1999 (30.12.99)</p> <p>(30) Priority Data: 09/224,426 31 December 1998 (31.12.98) US</p> <p>(63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application US 09/224,426 (CIP) Filed on 31 December 1998 (31.12.98)</p> <p>(71) Applicant (<i>for all designated States except US</i>): SYNAPTIC PHARMACEUTICAL CORPORATION [US/US]; 215 College Road, Paramus, NJ 07652 (US).</p> <p>(72) Inventors; and (75) Inventors/Applicants (<i>for US only</i>): SALON, John, A. [US/US]; Apartment A21, 180 Walnut Street, Montclair, NJ 07042 (US). LAZ, Thomas, M. [US/US]; 52 Fela Drive, Parlin, NJ 08859 (US). NAGORNY, Raisa [US/US]; 3-31 Cyril Avenue, Fair Lawn, NJ 07410 (US). WILSON, Amy, E. [US/US]; Apartment 2W, 310 East 92nd Street, New York, NY 10128 (US).</p>		<p>(74) Agent: WHITE, John, P.; Cooper &amp; Dunham LLP, 1185 Avenue of the Americas, New York, NY 10036 (US).</p> <p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p><b>Published</b> <i>Without international search report and to be republished upon receipt of that report.</i></p>	
<p>(54) Title: DNA ENCODING A HUMAN MELANIN CONCENTRATING HORMONE RECEPTOR (MCH1) AND USES THEREOF</p> <p>(57) Abstract</p> <p>This invention provides an isolated nucleic acid encoding a human MCH1 receptor, a purified human MCH1 receptor, vectors comprising isolated nucleic acid encoding a human MCH1 receptor, cells comprising such vectors, antibodies directed to a human MCH1 receptor, nucleic acid probes useful for detecting nucleic acid encoding human MCH1 receptors, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding human MCH1 receptors, transgenic, nonhuman animals which express DNA encoding a normal or mutant human MCH1 receptor, methods of isolating a human MCH1 receptor, methods of treating an abnormality that is linked to the activity of a human MCH1 receptor, as well as methods of determining binding of compounds to mammalian MCH1 receptors.</p>			

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DNA ENCODING A HUMAN MELANIN CONCENTRATING HORMONE RECEPTOR  
(MCH1) AND USES THEREOF

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BACKGROUND OF THE INVENTION

This application is a continuation-in-part of U.S. Serial No. 10 09/224,426, filed December 31, 1998, the contents of which is hereby incorporated by reference into the subject application.

Throughout this application, various publications are referenced in parentheses by author and year. Full citations for these references may be found at the end of the specification immediately preceding the sequence listings and the claims. The disclosure of these publications in their entireties are hereby incorporated by reference into this application to describe more fully the state of the art to 20 which this invention pertains.

Neuroregulators comprise a diverse group of natural products that subserve or modulate communication in the nervous system. They include, but are not limited to, 25 neuropeptides, amino acids, biogenic amines, lipids and lipid metabolites, and other metabolic byproducts. Many of these neuroregulator substances interact with specific cell surface receptors which transduce signals from the outside to the inside of the cell. G-protein coupled receptors (GPCRs) represent a major class of cell surface receptors with which many neurotransmitters interact to mediate their effects. GPCRs are predicted to have seven membrane- 30 spanning domains and are coupled to their effectors via G-proteins linking receptor activation with intracellular biochemical sequelae such as stimulation of adenylyl cyclase.

Melanin-concentrating hormone (MCH) is a cyclic peptide originally isolated from salmonid (teleost fish) pituitaries

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(Kawauchi et al., 1983). In fish the 17 amino acid peptide causes aggregation of melanin within the melanophores and inhibits the release of ACTH, acting as a functional antagonist of  $\alpha$ -MSH. Mammalian MCH (19 amino acids) is highly conserved between rat, mouse, and human, exhibiting 100% amino acid identity, but its physiological roles are less clear. MCH has been reported to participate in a variety of processes including feeding, water balance, energy metabolism, general arousal/attention state, memory and cognitive functions, and psychiatric disorders (for reviews, see Baker, 1991; Baker, 1994; Nahon, 1994; Knigge et al., 1996). Its role in feeding or body weight regulation is supported by a recent *Nature* publication (Qu et al., 1996) demonstrating that MCH is overexpressed in the hypothalamus of *ob/ob* mice compared with *ob/+* mice, and that fasting further increased MCH mRNA in both obese and normal mice during fasting. MCH also stimulated feeding in normal rats when injected into the lateral ventricles (Rossi et al., 1997). MCH also has been reported to functionally antagonize the behavioral effects of  $\alpha$ -MSH (Miller et al., 1993; Gonzalez et al., 1996; Sanchez et al., 1997); in addition, stress has been shown to increase POMC mRNA levels while decreasing the MCH precursor preproMCH (ppMCH) mRNA levels (Presse et al., 1992). Thus MCH may serve as an integrative neuropeptide involved in the reaction to stress, as well as in the regulation of feeding and sexual activity (Baker, 1991; Knigge et al., 1996).

The gene encoding the MCH precursor (ppMCH) has been cloned and encodes two additional peptides, neuropeptide EI (13 AA) and neuropeptide GE (19AA) (Nahon et al., 1989), which may also have biological activity. MCH peptide is synthesized primarily in hypothalamic neurons (the zona incerta and lateral hypothalamus) which project diffusely to many brain areas and to the pituitary (Bittencourt et al., 1992); NEI has also been identified in medium from explanted hypothalamic neurons (Parkes and Vale, 1993). Localization

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studies of the mRNA indicate that MCH is also present in the periphery (testes and GI tract; Hervieu and Nahon, 1995) but the highest concentrations are in the hypothalamus. There is also evidence for differential tissue-dependent processing of proMCH in mammals. A shorter MCH gene transcript that may result from alternate splicing was found in several brain areas and peripheral tissues, and a different peptide form was also found in the periphery (Viale et al., 1997). In humans, the gene encoding authentic MCH has been localized to chromosome 12, but two copies of a variant (truncated) gene are present on chromosome 5 (Breton et al., 1993); the functional significance, if any, of the variant is not yet known. Finally, the rat MCH gene may encode an additional putative peptide in a different reading frame (Toumaniantz et al., 1996).

Although the biological effects of MCH are believed to be mediated by specific receptors, binding sites for MCH have not been well described. A tritiated ligand ( $[^3\text{H}]\text{-MCH}$ ) was reported to exhibit specific binding to brain membranes but was unusable for saturation analyses, so neither affinity nor  $B_{\max}$  were determined (Drozdz and Eberle, 1995). Radioiodination of the tyrosine at position thirteen resulted in a ligand with dramatically reduced biological activity (see Drozdz and Eberle, 1995). In contrast, the radioiodination of the MCH analogue [ $\text{Phe}^{13}, \text{Tyr}^{19}$ ]-MCH was successful (Drozdz et al., 1995); the ligand retained biological activity and exhibited specific binding to a variety of cell lines including mouse melanoma (B16-F1, G4F, and G4F-7), PC12, and COS cells. In G4F-7 cells, the  $K_D = 0.118\text{nM}$  and the  $B_{\max} \sim 1100$  sites/cell. Importantly, the binding was not inhibited by  $\alpha$ -MSH but was weakly inhibited by rat ANF ( $K_i = 116\text{ nM}$  vs.  $12\text{ nM}$  for native MCH) (Drozdz et al., 1995). More recently specific MCH binding was reported in transformed keratinocytes (Burgaud et al., 1997) and melanoma cells (Drozdz et al., 1998), where photo-crosslinking studies suggest that the receptor is a membrane

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protein with an apparent molecular weight of 45-50 kDaltons, compatible with the molecular weight range of the GPCR superfamily of receptors. No radioautoradiographic studies of MCH receptor localization using this ligand have been reported as yet.

Signal transduction mechanisms for MCH receptors remain obscure. No direct evidence supporting G-protein coupling exists in mammals, but two lines of weak evidence exist in teleost fish for  $G_{\alpha q}$ - and/or  $G_{\alpha i}$ - type coupling: 1) indirect evidence exists for MCH acting via phospholipase C in teleost fish melanophores (phospholipase C inhibitors and protein kinase C inhibitors shift the MCH dose-response curve to the right, and TPA mimics MCH at low doses (Abrao et al., 1991)); and 2) MCH-elicited pigment aggregation in fish melanophores is associated with a reduction in basal cAMP levels, similar to that observed with norepinephrine (Svensson et al., 1991; Morishita et al., 1993). Arguing against G-protein coupling is the general structural homology of MCH with ANF, whose receptors are not in the GPCR superfamily. Recently the actions of MCH were reported to be mediated via activation of a phosphatidylinositol-3-kinase pathway which is typical of tyrosine kinase and cytokine receptors (Qu et al., 1998); however, since multiple signaling pathways (receptor cross talk) may produce this mediator no conclusions can be reached regarding MCH signal transduction pathways in mammalian systems.

The localization and biological activities of MCH peptide suggest that the modulation of MCH receptor activity may be useful in a number of therapeutic applications. The role of MCH in feeding is the best characterized of its potential clinical uses. MCH is expressed in the lateral hypothalamus, a brain area implicated in the regulation of thirst and hunger (Grillon et al., 1997); recently orexins A and B, which are potent orexigenic agents, have been shown to have very similar localization to MCH in the lateral

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hypothalamus (Sakurai et al., 1998). MCH mRNA levels in this brain region are increased in rats after 24 hours of food-deprivation (Hervé and Fellman, 1997); after insulin injection, a significant increase in the abundance and staining intensity of MCH immunoreactive perikarya and fibres was observed concurrent with a significant increase in the level of MCH mRNA (Bahjaoui-Bouhaddi et al., 1994). Consistent with the ability of MCH to stimulate feeding in rats (Rossi et al., 1997) is the observation that MCH mRNA levels are upregulated in the hypothalami of obese *ob/ob* mice (Qu et al., 1996), and decreased in the hypothalami of rats treated with leptin, whose food intake and body weight gains are also decreased (Sahu, 1998). MCH appears to act as a functional antagonist of the melanocortin system in its effects on food intake and on hormone secretion within the HPA (hypothalamopituitary/adrenal axis) (Ludwig et al., 1998). Together these data suggest a role for endogenous MCH in the regulation of energy balance and response to stress, and provide a rationale for the development of specific compounds acting at MCH receptors for use in the treatment of obesity and stress-related disorders.

In all species studied to date, a major portion of the neurons of the MCH cell group occupies a rather constant location in those areas of the lateral hypothalamus and subthalamus where they lie and may be a part of some of the so-called "extrapyramidal" motor circuits. These involve substantial striato- and pallidofugal pathways involving the thalamus and cerebral cortex, hypothalamic areas, and reciprocal connections to subthalamic nucleus, substantia nigra, and mid-brain centers (Bittencourt et al., 1992). In their location, the MCH cell group may offer a bridge or mechanism for expressing hypothalamic visceral activity with appropriate and coordinated motor activity. Clinically it may be of some value to consider the involvement of this MCH system in movement disorders, such as Parkinson's disease and Huntingdon's Chorea in which extrapyramidal circuits are known to be involved.

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Human genetic linkage studies have located authentic hMCH loci on chromosome 12 (12q23-24) and the variant hMCH loci on chromosome 5 (5q12-13) (Pedeutour et al., 1994). Locus 5 12q23-24 coincides with a locus to which autosomal dominant cerebellar ataxia type II (SCA2) has been mapped (Auburger et al., 1992; Twells et al., 1992). This disease comprises neurodegenerative disorders, including an olivopontocerebellar atrophy. Furthermore, the gene for 10 Darier's disease, has been mapped to locus 12q23-24 (Craddock et al., 1993). Darier's disease is characterized by abnormalities in keratinocyte adhesion and mental illnesses in some families. In view of the functional and 15 neuroanatomical patterns of the MCH neural system in the rat and human brains, the MCH gene may represent a good candidate for SCA2 or Darier's disease. Interestingly, diseases with high social impact have been mapped to this locus. Indeed, the gene responsible for chronic or acute forms of spinal muscular atrophies has been assigned to 20 chromosome 5q12-13 using genetic linkage analysis (Melki et al., 1990; Westbrook et al., 1992). Furthermore, independent lines of evidence support the assignment of a major schizophrenia locus to chromosome 5q11.2-13.3 (Sherrington et al., 1988; Bassett et al., 1988; Gilliam et 25 al., 1989). The above studies suggest that MCH may play a role in neurodegenerative diseases and disorders of emotion.

Additional therapeutic applications for MCH-related compounds are suggested by the observed effects of MCH in 30 other biological systems. For example, MCH may regulate reproductive functions in male and female rats. MCH transcripts and MCH peptide were found within germ cells in testes of adult rats, suggesting that MCH may participate in stem cell renewal and/or differentiation of early 35 spermatocytes (Hervieu et al., 1996). MCH injected directly into the medial preoptic area (MPOA) or ventromedial nucleus (VMN) stimulated sexual activity in female rats (Gonzalez et al., 1996). In ovariectomized rats primed with estradiol,

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MCH stimulated luteinizing hormone (LH) release while anti-MCH antiserum inhibited LH release (Gonzalez et al., 1997). The zona incerta, which contains a large population of MCH cell bodies, has previously been identified as a regulatory site for the pre-ovulatory LH surge (MacKenzie et al., 1984). MCH has been reported to influence release of pituitary hormones including ACTH and oxytocin. MCH analogues may also be useful in treating epilepsy. In the PTZ seizure model, injection of MCH prior to seizure induction prevented seizure activity in both rats and guinea pigs, suggesting that MCH-containing neurons may participate in the neural circuitry underlying PTZ-induced seizure (Knigge and Wagner, 1997). MCH has also been observed to affect behavioral correlates of cognitive functions. MCH treatment hastened extinction of the passive avoidance response in rats (McBride et al., 1994), raising the possibility that MCH receptor antagonists may be beneficial for memory storage and/or retention. A possible role for MCH in the modulation or perception of pain is supported by the dense innervation of the periaqueductal grey (PAG) by MCH-positive fibers. Finally, MCH may participate in the regulation of fluid intake. ICV infusion of MCH in conscious sheep produced diuretic, natriuretic, and kaliuretic changes in response to increased plasma volume (Parkes, 1996). Together with anatomical data reporting the presence of MCH in fluid regulatory areas of the brain, the results indicate that MCH may be an important peptide involved in the central control of fluid homeostasis in mammals.

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SUMMARY OF THE INVENTION

This invention provides an isolated nucleic acid encoding a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

This invention provides a nucleic acid encoding a human MCH1 receptor, wherein the nucleic acid (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when an MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

This invention provides a purified human MCH1 receptor protein.

This invention provides a vector comprising a nucleic acid encoding a human MCH1 receptor, particularly a vector adapted for expression of the human MCH1 receptor in mammalian or non-mammalian cells. One such vector is a plasmid designated pEXJ.HR-TL231 (ATCC Accession No. 203197) which comprises a nucleotide sequence encoding a human MCH1 receptor.

This invention also provides a cell comprising a vector which comprises a nucleic acid encoding a human MCH1 receptor as well as a membrane preparation isolated from such cells.

This invention further provides a nucleic acid probe comprising at least 15 nucleotides which specifically hybridizes with a nucleic acid encoding a mammalian MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within the nucleic acid which encodes the human MCH1 receptor or its complement,

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both of which are present in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197).

5 This invention further provides a nucleic acid probe comprising at least 15 nucleotides which specifically hybridizes with a nucleic acid encoding a mammalian MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within (a) the nucleic acid sequence shown in Figure 1 (SEQ ID NO: 1) or (b) the 10 reverse complement thereof.

15 This invention also provides an antisense oligonucleotide having a sequence capable of specifically hybridizing an RNA encoding a human MCH1 receptor, so as to prevent translation of the RNA and an antisense oligonucleotide having a sequence capable of specifically hybridizing to the genomic DNA encoding a human MCH1 receptor.

20 This invention further provides an antibody capable of binding to a human MCH1 receptor as well as an agent capable of competitively inhibiting the binding of the antibody to a human MCH1 receptor.

25 This invention provides a pharmaceutical composition comprising (a) an amount of the oligonucleotide described above capable of passing through a cell membrane and effective to reduce expression of a human MCH1 receptor and (b) a pharmaceutically acceptable carrier capable of passing through the cell membrane.

30 Moreover, this invention provides a transgenic, nonhuman mammal expressing DNA encoding a human MCH1 receptor. This invention also provides a transgenic, nonhuman mammal comprising a homologous recombination knockout of the native 35 human MCH1 receptor. This invention further provides a transgenic, nonhuman mammal whose genome comprises antisense DNA complementary to the DNA encoding a human MCH1 receptor so placed within the genome as to be transcribed into

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antisense mRNA which is complementary to mRNA encoding the human MCH1 receptor and which hybridizes to mRNA encoding the human MCH1 receptor, thereby reducing its translation.

5       In one embodiment this invention provides a process for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting cells containing DNA encoding and expressing on their cell surface a mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor.

15      This invention provides a process for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting a membrane preparation from cells transfected with DNA encoding and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor.

25      This invention provides a process involving competitive binding for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises separately contacting cells expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, a decrease in the binding of the second chemical compound to the mammalian MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the

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mammalian MCH1 receptor.

This invention provides a process involving competitive binding for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises separately contacting a membrane fraction from a cell extract of cells expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, a decrease in the binding of the second chemical compound to the mammalian MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the mammalian MCH1 receptor.

This invention provides a method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises (a) contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with a compound known to bind specifically to the mammalian MCH1 receptor; (b) contacting the preparation of step (a) with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor, under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor; (c) determining whether the binding of the compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so (d) separately determining the binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.

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This invention provides a method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises (a) contacting a membrane preparation from cells transfected with and expressing DNA encoding a mammalian MCH1 receptor with a compound known to bind specifically to the mammalian MCH1 receptor; (b) contacting the preparation of step (a) with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor, under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor; (c) determining whether the binding of the compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so (d) separately determining the binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.

This invention provides a method of detecting expression of a mammalian MCH1 receptor by detecting the presence of mRNA coding for the mammalian MCH1 receptor which comprises obtaining total mRNA from the cell and contacting the mRNA so obtained with a nucleic acid probe under hybridizing conditions, detecting the presence of mRNA hybridizing to the probe, and thereby detecting the expression of the mammalian MCH1 receptor by the cell.

This invention provides a method of detecting the presence of a mammalian MCH1 receptor on the surface of a cell which comprises contacting the cell with an antibody under conditions permitting binding of the antibody to the receptor, detecting the presence of the antibody bound to the cell, and thereby detecting the presence of the mammalian MCH1 receptor on the surface of the cell.

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This invention provides a method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a transgenic, nonhuman mammal whose levels of human MCH1 receptor activity are varied by use of an inducible promoter which regulates human MCH1 receptor expression.

This invention provides a method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a panel of transgenic, nonhuman mammals each expressing a different amount of human MCH1 receptor.

This invention provides a method for identifying an antagonist capable of alleviating an abnormality wherein the abnormality is alleviated by decreasing the activity of a human MCH1 receptor comprising administering a compound to the transgenic, nonhuman mammal and determining whether the compound alleviates the physical and behavioral abnormalities displayed by the transgenic, nonhuman mammal as a result of overactivity of a human MCH1 receptor, the alleviation of the abnormality identifying the compound as an antagonist. This invention also provides an antagonist identified by this method. This invention further provides a pharmaceutical composition comprising an antagonist identified by this method and a pharmaceutically acceptable carrier.

This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by decreasing the activity of a human MCH1 receptor which comprises administering to the subject an effective amount of this pharmaceutical composition, thereby treating the abnormality.

This invention provides a method for identifying an agonist capable of alleviating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of

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a human MCH1 receptor comprising administering a compound to a transgenic, nonhuman mammal, and determining whether the compound alleviates the physical and behavioral abnormalities displayed by the transgenic, nonhuman mammal,  
5 the alleviation of the abnormality identifying the compound as an agonist. This invention also provides an agonist identified by this method. This invention further provides a pharmaceutical composition comprising an agonist identified by this method and a pharmaceutically acceptable carrier. This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a human MCH1 receptor which comprises administering to the subject an effective amount of this pharmaceutical composition, thereby  
10 treating the abnormality.  
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This invention provides a method for diagnosing a predisposition to a disorder associated with the activity of a specific mammalian allele which comprises: (a) obtaining DNA of subjects suffering from the disorder; (b) performing a restriction digest of the DNA with a panel of restriction enzymes; (c) electrophoretically separating the resulting DNA fragments on a sizing gel; (d) contacting the resulting gel with a nucleic acid probe capable of specifically  
20 hybridizing with a unique sequence included within the sequence of a nucleic acid molecule encoding a human MCH1 receptor and labeled with a detectable marker; (e) detecting labeled bands which have hybridized to the DNA encoding a human MCH1 receptor labeled with a detectable marker to  
25 create a unique band pattern specific to the DNA of subjects suffering from the disorder; (f) preparing DNA obtained for diagnosis by steps (a)-(e); and (g) comparing the unique band pattern specific to the DNA of subjects suffering from the disorder from step (e) and the DNA obtained for  
30 diagnosis from step (f) to determine whether the patterns are the same or different and to diagnose thereby predisposition to the disorder if the patterns are the same.  
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This invention provides a method of preparing a purified human MCH1 receptor which comprises: (a) inducing cells to express the human MCH1 receptor; (b) recovering the human MCH1 receptor from the induced cells; and (c) purifying the human MCH1 receptor so recovered.

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This invention provides a method of preparing a purified human MCH1 receptor which comprises: (a) inserting nucleic acid encoding the human MCH1 receptor in a suitable vector; (b) introducing the resulting vector in a suitable host cell; (c) placing the resulting cell in suitable condition permitting the production of the isolated human MCH1 receptor; (d) recovering the human MCH1 receptor produced by the resulting cell; and (e) purifying the human MCH1 receptor so recovered.

This invention provides a process for determining whether a chemical compound is a mammalian MCH1 receptor agonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound under conditions permitting the activation of the mammalian MCH1 receptor, and detecting an increase in mammalian MCH1 receptor activity, so as to thereby determine whether the compound is a mammalian MCH1 receptor agonist. This invention also provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor agonist determined by this process effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a process for determining whether a chemical compound is a mammalian MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting the activation of the mammalian MCH1 receptor, and detecting a decrease in mammalian MCH1 receptor activity, so as to thereby determine

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whether the compound is a mammalian MCH1 receptor antagonist. This invention also provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor antagonist determined by this process effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a process for determining whether a chemical compound specifically binds to and activates a mammalian MCH1 receptor, which comprises contacting cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with the chemical compound under conditions suitable for activation of the mammalian MCH1 receptor, and measuring the second messenger response in the presence and in the absence of the chemical compound, a change in the second messenger response in the presence of the chemical compound indicating that the compound activates the mammalian MCH1 receptor.

This invention also provides a compound determined by this process. This invention further provides a pharmaceutical composition which comprises an amount of the compound (a MCH1 receptor agonist) determined by this process effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a process for determining whether a chemical compound specifically binds to and inhibits activation of a mammalian MCH1 receptor, which comprises separately contacting cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to activate the mammalian MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the mammalian MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical

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compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the mammalian MCH1 receptor. This invention also provides a compound determined by this process. This invention further provides a pharmaceutical composition which comprises an amount of the compound (a mammalian MCH1 receptor antagonist) determined by this effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a method of screening a plurality of chemical compounds not known to activate a mammalian MCH1 receptor to identify a compound which activates the mammalian MCH1 receptor which comprises: (a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds not known to activate the mammalian MCH1 receptor, under conditions permitting activation of the mammalian MCH1 receptor; (b) determining whether the activity of the mammalian MCH1 receptor is increased in the presence of the compounds; and if so (c) separately determining whether the activation of the mammalian MCH1 receptor is increased by each compound included in the plurality of compounds, so as to thereby identify the compound which activates the mammalian MCH1 receptor. This invention also provides a compound identified by this method. This invention further provides a pharmaceutical composition which comprises an amount of the compound (a mammalian MCH1 receptor agonist) identified by this method effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a method of screening a plurality of chemical compounds not known to inhibit the activation of a mammalian MCH1 receptor to identify a compound which

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inhibits the activation of the mammalian MCH1 receptor, which comprises: (a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting activation of the mammalian MCH1 receptor; (b) determining whether the activation of the mammalian MCH1 receptor is reduced in the presence of the plurality of compounds, relative to the activation of the mammalian MCH1 receptor in the absence of the plurality of compounds; and if so (c) separately determining the inhibition of activation of the mammalian MCH1 receptor for each compound included in the plurality of compounds, so as to thereby identify the compound which inhibits the activation of the mammalian MCH1 receptor.

This invention also provides a compound identified by this method. This invention further provides a pharmaceutical composition which comprises an amount of the compound (a mammalian MCH1 receptor antagonist) identified by this process effective to decrease activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.

This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor agonist effective to treat the abnormality.

This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by decreasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor antagonist effective to treat the abnormality.

This invention provides a process for making a composition of matter which specifically binds to a mammalian MCH1 receptor which comprises identifying a chemical compound

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using any of the processes described herein for identifying a compound which binds to and/or activates or inhibits activation of a mammalian MCH1 receptor and then synthesizing the chemical compound or a novel structural and functional analog or homolog thereof. This invention further provides a process for preparing a pharmaceutical composition which comprises administering a pharmaceutically acceptable carrier and a pharmaceutically acceptable amount of a chemical compound identified by any of the processes described herein for identifying a compound which binds to and/or activates or inhibits activation of a mammalian MCH1 receptor or a novel structural and functional analog or homolog thereof.

This invention provides a process for determining whether a chemical compound is a human MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH, and the cells do not express the MCH1 receptor prior to transfecting them.

This invention also provides a process for determining whether a chemical compound specifically binds to and inhibits activation of a human MCH1 receptor, which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the DNA encoding the human MCH1

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receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.

This invention further provides a method of screening a plurality of chemical compounds not known to inhibit the activation of a human MCH1 receptor to identify a compound which inhibits the activation of the human MCH1 receptor, which comprises:

- (a) contacting cells transfected with and expressing the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with the plurality of compounds in the presence of a known human MCH1 receptor agonist, under conditions permitting activation of the human MCH1 receptor, wherein the known MCH1 receptor agonist is MCH or a homolog or analog of MCH;
- (b) determining whether the activation of the human MCH1 receptor is reduced in the presence of the plurality of compounds, relative to the activation of the human

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MCH1 receptor in the absence of the plurality of compounds; and if so

- 5                         (c) separately determining the extent of inhibition of activation of the human MCH1 receptor for each compound included in the plurality of compounds, so as to thereby identify the compound which inhibits the activation of the human MCH1 receptor.
- 10                         This invention provides a process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.
- 15                         20                         30                         35                         This invention further provides a process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor

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and then synthesizing the chemical compound or a structural  
5 and functional analog or homolog thereof, wherein the  
chemical compound is identified as binding to the human MCH1  
receptor by a process involving competitive binding which  
comprises contacting a membrane preparation from cells  
expressing on their cell surface the human MCH1 receptor,  
with both the chemical compound and a second chemical  
compound known to bind to the receptor, and separately with  
only the second chemical compound, under conditions suitable  
10 for binding of both compounds, and detecting the extent of  
specific binding of the chemical compound to the human MCH1  
receptor, a decrease in the binding of the second chemical  
compound to the human MCH1 receptor in the presence of the  
chemical compound indicating that the chemical compound  
15 binds to the human MCH1 receptor, wherein the cells do not  
normally express the human MCH1 receptor, the human MCH1  
receptor is encoded by nucleic acid comprising the sequence  
shown in Figure 1 (Seq. ID No. 1) or contained in plasmid  
pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second  
20 chemical compound is MCH or a homolog or analog of MCH.

This invention also provides a process for making a  
composition of matter which is a human MCH1 receptor  
antagonist which comprises identifying a chemical compound  
25 which is a human MCH1 receptor antagonist and then  
synthesizing the chemical compound or a structural and  
functional analog or homolog thereof, wherein the chemical  
compound is identified as a human MCH1 receptor antagonist  
by a process which comprises contacting cells transfected  
30 with and expressing DNA encoding the human MCH1 receptor  
with the compound in the presence of a known human MCH1  
receptor agonist, under conditions permitting the activation  
of the human MCH1 receptor, and detecting a decrease in  
human MCH1 receptor activity, so as to thereby determine  
35 whether the compound is a human MCH1 receptor antagonist,  
wherein the cells do not normally express the human MCH1  
receptor, the human MCH1 receptor is encoded by nucleic acid  
comprising the sequence shown in Figure 1 (Seq. ID No. 1) or

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contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

- 5        This invention still further provides a process for making a composition of matter which specifically binds to and inhibits the activation of a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to and inhibits the activation of the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to and inhibiting the activation of the human MCH1 receptor by a process which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in 10      Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the 15      human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human 20      MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.
- 25      30      35      This invention provides a process for preparing a composition which comprises identifying a chemical compound which specifically binds to a human MCH1 receptor, and then

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admixing a carrier and the chemical compound or a structural  
and functional analog or homolog thereof, wherein the  
chemical compound is identified as binding to the human MCH1  
receptor by a process involving competitive binding which  
comprises contacting cells expressing on their cell surface  
the human MCH1 receptor, with both the chemical compound and  
a second chemical compound known to bind to the receptor,  
and separately with only the second chemical compound, under  
conditions suitable for binding of both compounds, and  
detecting the extent of specific binding of the chemical  
compound to the human MCH1 receptor, a decrease in the  
binding of the second chemical compound to the human MCH1  
receptor in the presence of the chemical compound indicating  
that the chemical compound binds to the human MCH1 receptor,  
wherein the cells do not normally express the human MCH1  
receptor, the human MCH1 receptor is encoded by nucleic acid  
comprising the sequence shown in Figure 1 (Seq. ID No. 1) or  
contained in plasmid pEXJ.HR-TL231 (ATCC Accession No.  
203197), and the second chemical compound is MCH or a  
homolog or analog of MCH.

This invention further provides a process for preparing a  
composition which comprises identifying a chemical compound  
which specifically binds to a human MCH1 receptor, and then  
admixing a carrier and the chemical compound or a structural  
and functional analog or homolog thereof, wherein the  
chemical compound is identified as binding to the human MCH1  
receptor by a process involving competitive binding which  
comprises contacting a membrane preparation from cells  
expressing on their cell surface the human MCH1 receptor,  
with both the chemical compound and a second chemical  
compound known to bind to the receptor, and separately with  
only the second chemical compound, under conditions suitable  
for binding of both compounds, and detecting the extent of  
specific binding of the chemical compound to the human MCH1  
receptor, a decrease in the binding of the second chemical  
compound to the human MCH1 receptor in the presence of the  
chemical compound indicating that the chemical compound

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5 binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

10 This invention also provides a process for preparing a composition which comprises identifying a chemical compound which is a human MCH1 receptor antagonist, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as a human MCH1 receptor antagonist  
15 by a process which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in  
20 human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or  
25 contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

30 This invention still further provides a process for preparing a composition which comprises identifying a chemical compound which specifically binds to and inhibits the activation of a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to and inhibiting activation of the human MCH1 receptor by a process which  
35 comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second

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messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.

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BRIEF DESCRIPTION OF THE FIGURES

Figure 1

Nucleotide sequence encoding a human MCH1 receptor (MCH1) (SEQ ID NO: 1). Three potential start (ATG) codons and the stop (TGA) codon are underlined.

Figure 2

Deduced amino acid sequence (SEQ ID NO: 2) of the human MCH1 receptor (MCH1) encoded by the nucleotide sequence shown Figure 1 (SEQ ID NO: 1).

Figure 3

Deduced amino acid sequence for human MCH1 (SEQ ID NO: 2). The seven putative transmembrane (TM) regions are underlined.

Figure 4

Nucleotide sequence of rat MCH1 (SEQ ID NO: 3). One start (ATG) codon and the stop codon (TGA) are underlined.

Figure 5

Deduced amino acid sequence for rat MCH1 (SEQ ID NO: 4).

Figure 6

MCH1-mediated PI dose response to MCH.

Figure 7

MCH1 challenge with several compounds of interest.

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Figure 8

MCH1-mediated extracellular acidification response to MCH and Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH. Results are reported as the average of two independent experiments performed in duplicate.

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Figure 9

Transcriptional response of MCH1-transfected Cos-7 cells to MCH.

5       Figure 10

Binding of [<sup>125</sup>I]Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH on MCH1-transfected Cos-7 cell membranes. Results are means ± S.E.M. (vertical lines) of triplicate determinations.

10      Figure 11

RT-PCR detection of MCH1 receptor mRNA in human mRNA samples.

Figure 12

15      Amino acid alignment of the N-terminal regions of MCH1 receptors encoded by various plasmids. The mutations present in R106 (SEQ ID NO: 16) and R114 (SEQ ID NO: 17) are shown in lower case. Potential initiating methionines are shown in bold. The amino acid sequence downstream of position 100  
20      is identical for all four plasmids.

Figure 13

Amino acid sequence (SEQ ID NO: 27) of the mutant human MCH1 receptor encoded by plasmid R106.

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Figure 14

Amino acid sequence (SEQ ID NO: 28) of the mutant human MCH1 receptor encoded by plasmid R114.

30      Figure 15

Amino acid sequence (SEQ ID NO: 29) of the mutant human MCH1 receptor encoded by plasmid B0120.

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DETAILED DESCRIPTION OF THE INVENTION

Throughout this application, the following standard abbreviations are used to indicate specific nucleotide bases:

	A =	adenine
	G =	guanine
	C =	cytosine
5	T =	thymine
	U =	uracil
	M =	adenine or cytosine
10	R =	adenine or guanine
	W =	adenine, thymine, or uracil
15	S =	cytosine or guanine
	Y =	cytosine, thymine, or uracil
	K =	guanine, thymine, or uracil
	V =	adenine, cytosine, or guanine (not thymine or uracil)
20	H =	adenine, cytosine, thymine, or uracil (not guanine)
	D =	adenine, guanine, thymine, or uracil (not cytosine)
	B =	cytosine, guanine, thymine, or uracil (not adenine)
25	N =	adenine, cytosine, guanine, thymine, or uracil (or other modified base such as inosine)
	I =	inosine

Furthermore, the term "agonist" is used throughout this application to indicate any peptide or non-peptidyl compound which increases the activity of any of the polypeptides of the subject invention. The term "antagonist" is used throughout this application to indicate any peptide or non-peptidyl compound which decreases the activity of any of the polypeptides of the subject invention. The term "mammalian" is used throughout this invention to include mutant forms of the human MCH1 receptor.

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The activity of a G-protein coupled receptor such as the polypeptides disclosed herein may be measured using any of a variety of functional assays in which activation of the receptor in question results in an observable change in the level of some second messenger system, including, but not limited to, adenylate cyclase, calcium mobilization, arachidonic acid release, ion channel activity, inositol phospholipid hydrolysis or guanylyl cyclase. Heterologous expression systems utilizing appropriate host cells to express the nucleic acid of the subject invention are used to obtain the desired second messenger coupling. Receptor activity may also be assayed in an oocyte expression system.

It is possible that the human MCH1 receptor gene contains introns and furthermore, the possibility exists that additional introns could exist in coding or non-coding regions. In addition, spliced form(s) of mRNA may encode additional amino acids either upstream of the currently defined starting methionine or within the coding region. Further, the existence and use of alternative exons is possible, whereby the mRNA may encode different amino acids within the region comprising the exon. In addition, single amino acid substitutions may arise via the mechanism of RNA editing such that the amino acid sequence of the expressed protein is different than that encoded by the original gene. (Burns et al., 1996; Chu et al., 1996). Such variants may exhibit pharmacologic properties differing from the polypeptide encoded by the original gene.

This invention provides splice variants of the human MCH1 receptor disclosed herein. This invention further provides for alternate translation initiation sites and alternately spliced or edited variants of nucleic acids encoding the human MCH1 receptor of this invention.

The nucleic acid of the subject invention also includes nucleic acid analogs of the human MCH1 receptor gene, wherein the human MCH1 receptor gene comprises the nucleic

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acid sequence shown in Fig. 1 or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197). Nucleic acid analogs of the human MCH1 receptor genes differ from the human MCH1 receptor gene described herein in terms of the identity or location of one or more nucleic acid bases (deletion analogs containing less than all of the nucleic acid bases shown in Fig. 1 or contained in plasmid pEXJ.HR-TL231, substitution analogs wherein one or more nucleic acid bases shown in Fig. 1 or contained in plasmids pEXJ.HR-TL231 are replaced by other nucleic acid bases, and addition analogs, wherein one or more nucleic acid bases are added to a terminal or medial portion of the nucleic acid sequence) and which encode proteins which share some or all of the properties of the proteins encoded by the nucleic acid sequences shown in Fig. 1 or contained in plasmid pEXJ.HR-TL231. In one embodiment of the present invention, the nucleic acid analog encodes a protein which comprises an amino acid sequence as shown in Fig. 2 or encoded by the nucleic acid sequence contained in plasmid pEXJ.HR-TL231. In another embodiment, the nucleic acid analog encodes a protein comprising an amino acid sequence which differs from the amino acid sequences shown in Fig. 2 or encoded by the nucleic acid contained in plasmids pEXJ.HR-TL231. In a further embodiment, the protein encoded by the nucleic acid analog has a function which is the same as the function of the receptor protein comprising the amino acid sequence shown in Fig. 2. In another embodiment, the function of the protein encoded by the nucleic acid analog differs from the function of the receptor protein comprising the amino acid sequence shown in Fig. 2. In another embodiment, the variation in the nucleic acid sequence occurs within the transmembrane (TM) region of the protein. In a further embodiment, the variation in the nucleic acid sequence occurs outside of the TM region.

This invention provides the above-described isolated nucleic acid, wherein the nucleic acid is DNA. In an embodiment, the DNA is cDNA. In another embodiment, the DNA is genomic

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DNA. In still another embodiment, the nucleic acid is RNA. Methods for production and manipulation of nucleic acid molecules are well known in the art.

5 This invention further provides nucleic acid which is degenerate with respect to the DNA encoding the polypeptides described herein. In an embodiment, the nucleic acid comprises a nucleotide sequence which is degenerate with respect to the nucleotides sequence shown in Figure 1 (SEQ ID NO: 2) or the nucleotide sequence contained in the 10 plasmid pEXJ.HR-TL231, that is, a nucleotide sequence which is translated into the same amino acid sequence.

15 This invention also encompasses DNAs and cDNAs which encode amino acid sequences which differ from those of the polypeptides of this invention, but which should not produce phenotypic changes. Alternately, this invention also encompasses DNAs, cDNAs, and RNAs which hybridize to the 20 DNA, cDNA, and RNA of the subject invention. Hybridization methods are well known to those of skill in the art.

The nucleic acids of the subject invention also include 25 nucleic acid molecules coding for polypeptide analogs, fragments or derivatives of antigenic polypeptides which differ from naturally-occurring forms in terms of the identity or location of one or more amino acid residues (deletion analogs containing less than all of the residues specified for the protein, substitution analogs wherein one or more residues specified are replaced by other residues 30 and addition analogs wherein one or more amino acid residues is added to a terminal or medial portion of the polypeptides) and which share some or all properties of naturally-occurring forms. These molecules include: the incorporation of codons "preferred" for expression by 35 selected non-mammalian hosts; the provision of sites for cleavage by restriction endonuclease enzymes; and the provision of additional initial, terminal or intermediate DNA sequences that facilitate construction of readily

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expressed vectors. The creation of polypeptide analogs is well known to those of skill in the art (R.F. Spurney et al. (1997); Fong, T.M. et al. (1995); Underwood, D.J. et al. (1994); Graziano, M.P. et al. (1996); Guan X.M. et al. (1995)).

The modified polypeptides of this invention may be transfected into cells either transiently or stably using methods well-known in the art, examples of which are disclosed herein. This invention also provides for binding assays using the modified polypeptides, in which the polypeptide is expressed either transiently or in stable cell lines. This invention further provides a compound identified using a modified polypeptide in a binding assay such as the binding assays described herein.

The nucleic acids described and claimed herein are useful for the information which they provide concerning the amino acid sequence of the polypeptide and as products for the large scale synthesis of the polypeptides by a variety of recombinant techniques. The nucleic acid molecule is useful for generating new cloning and expression vectors, transformed and transfected prokaryotic and eukaryotic host cells, and new and useful methods for cultured growth of such host cells capable of expression of the polypeptide and related products.

This invention provides an isolated nucleic acid encoding a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In one embodiment, the nucleic acid is DNA. In another embodiment, the DNA is cDNA. In another embodiment, the DNA is genomic DNA. In another embodiment, the nucleic acid is RNA.

This invention also provides methods of using an isolated nucleic acid encoding species homologs of the MCH1 receptor encoded by the nucleic acid sequence shown in Fig. 1 (SEQ ID

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- NO: 1) or encoded by the plasmid pEXJ.HR-TL231. In one embodiment, the nucleic acid encodes a mammalian MCH1 receptor homolog which has substantially the same amino acid sequence as does the MCH1 receptor encoded by the plasmid pEXJ.HR-TL231. In another embodiment, the nucleic acid encodes a mammalian MCH1 receptor homolog which has above 5 65% amino acid identity to the MCH1 receptor encoded by the plasmid pEXJ.HR-TL231; preferably above 75% amino acid identity to the MCH1 receptor encoded by the plasmid pEXJ.HR-TL231; more preferably above 85% amino acid identity to the MCH1 receptor encoded by the plasmid pEXJ.HR-TL231; most preferably above 95% amino acid identity to the MCH1 receptor encoded by the plasmid pEXJ.HR-TL231. In another embodiment, the mammalian MCH1 receptor homolog has above 10 70% nucleic acid identity to the MCH1 receptor gene contained in plasmid pEXJ.HR-TL231; preferably above 80% nucleic acid identity to the MCH1 receptor gene contained in the plasmid pEXJ.HR-TL231; more preferably above 90% nucleic acid identity to the MCH1 receptor gene contained in the 15 plasmid pEXJ.HR-TL231. Examples of methods for isolating and purifying species homologs are described elsewhere (e.g., U.S. Patent No. 5,602,024, WO94/14957, WO97/26853, 20 WO98/15570).
- In a separate embodiment of the present invention, the 25 nucleic acid encodes a MCH1 receptor which has an amino acid sequence identical to that encoded by the plasmid pEXJ.HR-TL231. In a further embodiment, the MCH1 receptor comprises a sequence substantially the same as the amino acid sequence shown in Figure 2 (SEQ ID NO: 2). In another embodiment, 30 the MCH1 receptor comprises an amino acid sequence as shown in Figure 2 (SEQ ID NO: 2).
- In one embodiment, the mutant human MCH1 receptor comprises 35 an amino acid sequence as shown in Figure 13 (SEQ ID NO: 27). In another embodiment, the mutant human MCH1 receptor comprises an amino acid sequence as shown in Figure 14 (SEQ ID NO: 28). In still another embodiment, the mutant human

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MCH1 receptor comprises an amino acid sequence as shown in Figure 15 (SEQ ID NO: 29).

5 In separate embodiments, the human MCH1 receptor is encoded by the nucleic acid sequence shown in Figure 1 beginning with any of the three indicated start (ATG) codons.

10 This invention provides an isolated nucleic acid encoding a modified human MCH1 receptor, which differs from a human MCH1 receptor by having an amino acid(s) deletion, replacement, or addition in the third intracellular domain.

15 This invention provides a nucleic acid encoding a human MCH1 receptor, wherein the nucleic acid (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

20 Hybridization at low stringency is performed at 40°C in a hybridization buffer containing 25% formamide, 5X SCC, 7mM Tris, 1X Denhardt's, 25μl/ml salmon sperm DNA. Wash at 40°C in 0.1X SCC, 0.1% SDS. Changes in pH are measured through microphysiometric measurement of receptor mediated extracellular acidification rates. Because cellular metabolism is intricately involved in a broad range of cellular events (including receptor activation of multiple messenger pathways), the use of microphysiometric measurements of cell metabolism can in principle provide a generic assay of cellular activity arising from the activation of any receptor regardless of the specifics of the receptor's signaling pathway. General guidelines for transient receptor expression, cell preparation and microphysiometric recording are described elsewhere (Salon, J.A. and Owicki, J.A., 1996). Receptors and/or control vectors are transiently expressed in CHO-K1 cells, by

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liposome mediated transfection according to the manufacturers recommendations (LipofectAMINE, GibcoBRL, Gaithersburg, MD), and maintained in Ham's F-12 complete (10% serum). A total of 10 $\mu$ g of DNA is used to transfect each 75cm<sup>2</sup> flask which had been split 24 hours prior to the transfection and judged to be 70-80% confluent at the time of transfection. 24 hours post transfection, the cells are harvested and 3 x 10<sup>5</sup> cells seeded into microphysiometer capsules. Cells are allowed to attach to the capsule membrane for an additional 24 hours; during the last 16 hours, the cells are switched to serum-free F-12 complete to minimize ill-defined metabolic stimulation caused by assorted serum factors. On the day of the experiment the cell capsules are transferred to the microphysiometer and allowed to equilibrate in recording media (low buffer RPMI 1640, no bicarbonate, no serum (Molecular Devices Corporation, Sunnyvale, CA) containing 0.1% fatty acid free BSA), during which a baseline measurement of basal metabolic activity is established. A standard recording protocol specifies a 100 $\mu$ l/min flow rate, with a 2 min total pump cycle which includes a 30 sec flow interruption during which the acidification rate measurement is taken. Ligand challenges involve a 1 min 20 sec exposure to the sample just prior to the first post challenge rate measurement being taken, followed by two additional pump cycles for a total of 5 min 20 sec sample exposure. Typically, drugs in a primary screen are presented to the cells at 10 $\mu$ M final concentration. Ligand samples are then washed out and the acidification rates reported are expressed as a percentage increase of the peak response over the baseline rate observed just prior to challenge. An examples of a MCH ligand includes, but is not limited to, the endogenous MCH peptide.

This invention provides a purified human MCH1 receptor protein.

This invention provides a vector comprising nucleic acid

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encoding a human MCH1 receptor. In an embodiment, the vector is adapted for expression in a cell which comprises the regulatory elements necessary for expression of the nucleic acid in the cell operatively linked to the nucleic acid encoding the human MCH1 receptor as to permit expression thereof. In separate embodiments, the cell is a bacterial cell, an amphibian cell, a yeast cell, an insect cell or a mammalian cell. In another embodiment, the vector is a baculovirus. In one embodiment, the vector is a plasmid.

This invention provides a plasmid designated pEXJ.HR-TL231 (ATCC Accession No. 203197). This plasmid comprises the regulatory elements necessary for expression of DNA in a mammalian cell operatively linked to DNA encoding the human MCH1 receptor so as to permit expression thereof.

This plasmid (pEXJ.HR-TL231) was deposited on September 17, 1998, with the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, Maryland 20852, U.S.A. under the provisions of the Budapest Treaty for the International Recognition of the Deposit of Microorganisms for the Purposes of Patent Procedure and was accorded ATCC Accession No. 203197.

This invention further provides for any vector or plasmid which comprises modified untranslated sequences, which are beneficial for expression in desired host cells or for use in binding or functional assays. For example, a vector or plasmid with untranslated sequences of varying lengths may express differing amounts of the polypeptide depending upon the host cell used. In an embodiment, the vector or plasmid comprises the coding sequence of the polypeptide and the regulatory elements necessary for expression in the host cell.

This invention provides a cell comprising a vector comprising a nucleic acid encoding the human MCH1 receptor.

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In an embodiment, the cell is a non-mammalian cell. In a further embodiment, the non-mammalian cell is a *Xenopus* oocyte cell or a *Xenopus melanophore* cell. In another embodiment, the cell is a mammalian cell. In a further embodiment, the mammalian cell is a COS-7 cell, a 293 human embryonic kidney cell, a NIH-3T3 cell, a LM(tk-) cell, a mouse Y1 cell, or a CHO cell.

This invention provides an insect cell comprising a vector adapted for expression in an insect cell which comprises a nucleic acid encoding a human MCH1 receptor. In another embodiment, the insect cell is an Sf9 cell, an Sf21 cell or a *Trichoplusia ni* 5B1-4 (HighFive) cell.

This invention provides a membrane preparation isolated from any one of the cells described above.

This invention provides a nucleic acid probe comprising at least 15 nucleotides, which probe specifically hybridizes with a nucleic acid encoding a human MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within one of the two strands of the nucleic acid encoding a human MCH1 receptor present in plasmid pEXJ.HR-TL231. This invention also provides a nucleic acid probe comprising at least 15 nucleotides, which probe specifically hybridizes with a nucleic acid encoding a human MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within (a) the nucleic acid sequence shown in Figure 1 (SEQ ID NO: 1) or (b) the reverse complement thereto. In one embodiment, the nucleic acid is DNA. In another embodiment, the nucleic acid is RNA.

As used herein, the phrase "specifically hybridizing" means the ability of a nucleic acid molecule to recognize a nucleic acid sequence complementary to its own and to form double-helical segments through hydrogen bonding between complementary base pairs.

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Nucleic acid probe technology is well known to those skilled in the art who will readily appreciate that such probes may vary greatly in length and may be labeled with a detectable label, such as a radioisotope or fluorescent dye, to facilitate detection of the probe. DNA probe molecules may be produced by insertion of a DNA molecule which encodes the polypeptides of this invention into suitable vectors, such as plasmids or bacteriophages, followed by transforming into suitable bacterial host cells, replication in the transformed bacterial host cells and harvesting of the DNA probes, using methods well known in the art. Alternatively, probes may be generated chemically from DNA synthesizers.

RNA probes may be generated by inserting the DNA molecule which encodes the polypeptides of this invention downstream of a bacteriophage promoter such as T3, T7, or SP6. Large amounts of RNA probe may be produced by incubating the labeled nucleotides with the linearized fragment where it contains an upstream promoter in the presence of the appropriate RNA polymerase.

This invention provides an antisense oligonucleotide having a sequence capable of specifically hybridizing to RNA encoding a human MCH1 receptor, so as to prevent translation of the RNA. This invention also provides an antisense oligonucleotide having a sequence capable of specifically hybridizing to genomic DNA encoding a human MCH1 receptor. In one embodiment, the oligonucleotide comprises chemically modified nucleotides or nucleotide analogues.

This invention provides an antibody capable of binding to a human MCH1 receptor encoded by a nucleic acid encoding a human MCH1 receptor. This invention also provides an agent capable of competitively inhibiting the binding of the antibody to a human MCH1 receptor. In one embodiment, the antibody is a monoclonal antibody or antisera.

This invention provides a pharmaceutical composition

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comprising (a) an amount of the oligonucleotide capable of passing through a cell membrane and effective to reduce expression of a human MCH1 receptor and (b) a pharmaceutically acceptable carrier capable of passing through the cell membrane. In an embodiment, the oligonucleotide is coupled to a substance which inactivates mRNA. In a further embodiment, the substance which inactivates mRNA is a ribozyme. In another embodiment, the pharmaceutically acceptable carrier comprises a structure which binds to a human MCH1 receptor on a cell capable of being taken up by the cells after binding to the structure. In a further embodiment, the pharmaceutically acceptable carrier is capable of binding to a human MCH1 receptor which is specific for a selected cell type.

This invention provides a pharmaceutical composition which comprises an amount of an antibody effective to block binding of a ligand to a human MCH1 receptor and a pharmaceutically acceptable carrier.

As used herein, the phrase "pharmaceutically acceptable carrier" means any of the standard pharmaceutically acceptable carriers. Examples include, but are not limited to, phosphate buffered saline, physiological saline, water, and emulsions, such as oil/water emulsions.

This invention provides a transgenic, nonhuman mammal expressing DNA encoding a human MCH1 receptor. This invention also provides a transgenic, nonhuman mammal comprising a homologous recombination knockout of the native human MCH1 receptor. This invention further provides a transgenic, nonhuman mammal whose genome comprises antisense DNA complementary to the DNA encoding a human MCH1 receptor so placed within the genome as to be transcribed into antisense mRNA which is complementary to mRNA encoding the human MCH1 receptor and which hybridizes to mRNA encoding the human MCH1 receptor, thereby reducing its translation. In an embodiment, the DNA encoding the human MCH1 receptor

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additionally comprises an inducible promoter. In another embodiment, the DNA encoding the human MCH1 receptor additionally comprises tissue specific regulatory elements. In a further embodiment, the transgenic, nonhuman mammal is  
5 a mouse.

Animal model systems which elucidate the physiological and behavioral roles of the polypeptides of this invention are produced by creating transgenic animals in which the  
10 activity of the polypeptide is either increased or decreased, or the amino acid sequence of the expressed polypeptide is altered, by a variety of techniques. Examples of these techniques include, but are not limited to: 1) Insertion of normal or mutant versions of DNA  
15 encoding the polypeptide, by microinjection, electroporation, retroviral transfection or other means well known to those in the art, into appropriate fertilized embryos in order to produce a transgenic animal or 2) Homologous recombination of mutant or normal, human or  
20 animal versions of these genes with the native gene locus in transgenic animals to alter the regulation of expression or the structure of these polypeptide sequences. The technique of homologous recombination is well known in the art. It replaces the native gene with the inserted gene and so is useful for producing an animal that cannot express native  
25 polypeptides but does express, for example, an inserted mutant polypeptide, which has replaced the native polypeptide in the animal's genome by recombination, resulting in underexpression of the transporter.  
30 Microinjection adds genes to the genome, but does not remove them, and so is useful for producing an animal which expresses its own and added polypeptides, resulting in overexpression of the polypeptides.

35 One means available for producing a transgenic animal, with a mouse as an example, is as follows: Female mice are mated, and the resulting fertilized eggs are dissected out of their oviducts. The eggs are stored in an appropriate medium such

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as M2 medium. DNA or cDNA encoding a polypeptide of this invention is purified from a vector by methods well known in the art. Inducible promoters may be fused with the coding region of the DNA to provide an experimental means to regulate expression of the trans-gene. Alternatively, or in addition, tissue specific regulatory elements may be fused with the coding region to permit tissue-specific expression of the trans-gene. The DNA, in an appropriately buffered solution, is put into a microinjection needle (which may be made from capillary tubing using a pipette puller) and the egg to be injected is put in a depression slide. The needle is inserted into the pronucleus of the egg, and the DNA solution is injected. The injected egg is then transferred into the oviduct of a pseudopregnant mouse ( a mouse stimulated by the appropriate hormones to maintain pregnancy but which is not actually pregnant ), where it proceeds to the uterus, implants, and develops to term. As noted above, microinjection is not the only method for inserting DNA into the egg cell, and is used here only for exemplary purposes.

This invention provides a process for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting cells comprising DNA encoding, and expressing on their cell surface, the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement. This invention also provides a process for identifying a chemical compound which

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specifically binds to a mammalian MCH1 receptor which comprises contacting a membrane preparation from cells comprising DNA encoding, and expressing on their cell surface, the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement. In one embodiment, the MCH1 receptor is a human MCH1 receptor. In another embodiment, the MCH1 receptor is a rat MCH1 receptor. In another embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid sequence as the sequence of the human MCH1 receptor encoded by plasmid pEXJ.HR-TL231. In a further embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid sequence as that shown in Figure 2 (SEQ ID NO: 2). In another embodiment, the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 2 (SEQ ID NO: 2). In a different embodiment, the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 13 (SEQ ID NO: 27). In another embodiment, the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 14 (SEQ ID NO: 28). In still another embodiment, the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 15 (SEQ ID NO: 29). In one embodiment, the compound is not previously known to bind to a mammalian MCH1 receptor. This invention further provides a compound identified by the above-described processes.

In one embodiment of the above-described processes, the cell

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is an insect cell. In another embodiment, the cell is a mammalian cell. In a further embodiment, the cell is nonneuronal in origin. In a further embodiment, the nonneuronal cell is a COS-7 cell, 293 human embryonic kidney cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.

This invention provides a process involving competitive binding for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting cells expressing on their cell surface the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, a decrease in the binding of the second chemical compound to the mammalian MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

This invention also provides a process involving competitive binding for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting a membrane preparation from cells expressing on their cell surface the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and

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separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, a decrease in the binding of the 5 second chemical compound to the mammalian MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor 10 (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added 15 to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

In one embodiment, the mammalian MCH1 receptor is a human 20 MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor. In another embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid 25 sequence as the human MCH1 receptor encoded by plasmid pEXJ.HR-TL231. In a further embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid sequence as that shown in Figure 2 (SEQ ID NO: 2). In another embodiment, the mammalian MCH1 receptor comprises 30 the amino acid sequence shown in Figure 2 (SEQ ID NO: 2).

In one embodiment, the cell is an insect cell. In another embodiment, the cell is a mammalian cell. In a further embodiment, the cell is nonneuronal in origin. In another 35 embodiment, the nonneuronal cell is a COS-7 cell, 293 human embryonic kidney cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell. In one embodiment, the compound is not previously known to bind to a mammalian MCH1

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receptor.

This invention provides a compound identified by the above-described processes.

- 5        This invention provides a method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises (a) contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor, under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor; (b) determining whether the binding of a compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so (c) separately determining the binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.
- 25      This invention provides a method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises (a) contacting a membrane preparation from cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor, under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor; (b) determining whether the binding of a compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so (c) separately determining the
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binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.

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In one embodiment of the above-described methods, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor. In another embodiment, the cell is a mammalian cell. In a further embodiment, the mammalian cell is non-neuronal in origin. In another embodiment, the non-neuronal cell is a COS-7 cell, a 293 human embryonic kidney cell, a LM(tk-) cell, a CHO cell, a mouse Y1 cell, or an NIH-3T3 cell.

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This invention also provides a method of detecting expression of a mammalian MCH1 receptor by detecting the presence of mRNA coding for the mammalian MCH1 receptor which comprises obtaining total mRNA from the cell and contacting the mRNA so obtained from a nucleic acid probe under hybridizing conditions, detecting the presence of mRNA hybridizing to the probe, and thereby detecting the expression of the mammalian MCH1 receptor by the cell.

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This invention further provides a method of detecting the presence of a mammalian MCH1 receptor on the surface of a cell which comprises contacting the cell with an antibody under conditions permitting binding of the antibody to the receptor, detecting the presence of the antibody bound to the cell, and thereby detecting the presence of the mammalian MCH1 receptor on the surface of the cell.

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This invention provides a method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a transgenic, nonhuman mammal whose levels of human MCH1 receptor activity are varied by use of an inducible promoter which regulates

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human MCH1 receptor expression.

This invention also provides a method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a panel of transgenic, nonhuman mammals each expressing a different amount of human MCH1 receptor.

This invention provides a method for identifying an antagonist capable of alleviating an abnormality wherein the abnormality is alleviated by decreasing the activity of a human MCH1 receptor comprising administering a compound to a transgenic, nonhuman mammal, and determining whether the compound alleviates the physical and behavioral abnormalities displayed by the transgenic, nonhuman mammal as a result of overactivity of a human MCH1 receptor, the alleviation of the abnormality identifying the compound as an antagonist. This invention also provides an antagonist identified by the above-described method. This invention further provides a pharmaceutical composition comprising an antagonist identified by the above-described method and a pharmaceutically acceptable carrier. This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by decreasing the activity of a human MCH1 receptor which comprises administering to the subject an effective amount of this pharmaceutical composition, thereby treating the abnormality.

This invention provides a method for identifying an agonist capable of alleviating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a human MCH1 receptor comprising administering a compound to transgenic, nonhuman mammal, and determining whether the compound alleviates the physical and behavioral abnormalities displayed by the transgenic, nonhuman mammal, the alleviation of the abnormality identifying the compound as an agonist. This invention also provides an agonist

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identified by the above-described method. This invention further provides a pharmaceutical composition comprising an agonist identified by the above-described method and a pharmaceutically acceptable carrier. This invention further provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a human MCH1 receptor which comprises administering to the subject an effective amount of this pharmaceutical composition, thereby treating the abnormality.

This invention provides a method for diagnosing a predisposition to a disorder associated with the activity of a specific mammalian allele which comprises: (a) obtaining DNA of subjects suffering from the disorder; (b) performing a restriction digest of the DNA with a panel of restriction enzymes; (c) electrophoretically separating the resulting DNA fragments on a sizing gel; (d) contacting the resulting gel with a nucleic acid probe capable of specifically hybridizing with a unique sequence included within the sequence of a nucleic acid molecule encoding a human MCH1 receptor and labeled with a detectable marker; (e) detecting labeled bands which have hybridized to the DNA encoding a human MCH1 receptor labeled with a detectable marker to create a unique band pattern specific to the DNA of subjects suffering from the disorder; (f) preparing DNA obtained for diagnosis by steps (a)-(e); and (g) comparing the unique band pattern specific to the DNA of subjects suffering from the disorder from step (e) and the DNA obtained for diagnosis from step (f) to determine whether the patterns are the same or different and to diagnose thereby predisposition to the disorder if the patterns are the same. In one embodiment, a disorder associated with the activity of a specific mammalian allele is diagnosed.

This invention provides a method of preparing the purified human MCH1 receptor which comprises: (a) inducing cells to express the human MCH1 receptor; (b) recovering the human

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MCH1 receptor from the induced cells; and (c) purifying the human MCH1 receptor so recovered.

This invention provides a method of preparing the purified human MCH1 receptor which comprises: (a) inserting nucleic acid encoding the human MCH1 receptor in a suitable vector; (b) introducing the resulting vector in a suitable host cell; (c) placing the resulting cell in suitable condition permitting the production of the isolated human MCH1 receptor; (d) recovering the human MCH1 receptor produced by the resulting cell; and (e) purifying the human MCH1 receptor so recovered.

This invention provides a process for determining whether a chemical compound is a mammalian MCH1 receptor agonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound under conditions permitting the activation of the mammalian MCH1 receptor, and detecting an increase in mammalian MCH1 receptor activity, so as to thereby determine whether the compound is a mammalian MCH1 receptor agonist. This invention also provides a process for determining whether a chemical compound is a mammalian MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting the activation of the mammalian MCH1 receptor, and detecting a decrease in mammalian MCH1 receptor activity, so as to thereby determine whether the compound is a mammalian MCH1 receptor antagonist. In one embodiment, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

This invention further provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor agonist determined by the above-described process effective

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to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier. In one embodiment, the mammalian MCH1 receptor agonist is not previously known.

5       This invention provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor antagonist determined by the above-described process effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier. In one embodiment, the mammalian MCH1 receptor antagonist is not previously known.

10      This invention provides a process for determining whether a chemical compound specifically binds to and activates a mammalian MCH1 receptor, which comprises contacting cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with the chemical compound under conditions suitable for activation of the mammalian MCH1 receptor, and measuring the second messenger response in the presence and in the absence 15      of the chemical compound, a change in the second messenger response in the presence of the chemical compound indicating that the compound activates the mammalian MCH1 receptor. In one embodiment, the second messenger response comprises chloride channel activation and the change in second messenger is an increase in the level of inward chloride current.

20      This invention also provides a process for determining whether a chemical compound specifically binds to and inhibits activation of a mammalian MCH1 receptor, which comprises separately contacting cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to activate 25      the mammalian MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation

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of the mammalian MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the mammalian MCH1 receptor. In one embodiment, the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound. This invention also provides the above-described processes performed with membrane preparations from cells producing a second messenger response and transfected with and expressing the mammalian MCH1 receptor.

In one embodiment of the above-described processes, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor. In another embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid sequence as encoded by the plasmid pEXJ.HR-TL231. In a further embodiment, the mammalian MCH1 receptor comprises substantially the same amino acid sequence as that shown in Figure 2 (SEQ ID NO: 2). In another embodiment, the mammalian MCH1 receptor comprises an amino acid sequence as shown in Figure 2 (SEQ ID NO: 2). In an embodiment, the cell is an insect cell. In a further embodiment, the cell is a mammalian cell. In a still further embodiment, the mammalian cell is nonneuronal in origin. In another embodiment, the nonneuronal cell is a COS-7 cell, CHO cell, 293 human embryonic kidney cell, NIH-3T3 cell or LM(tk-) cell. In an embodiment, the compound is not previously known to bind to

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a mammalian MCH1 receptor. This invention also provides a compound determined by the above-described processes.

This invention also provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor agonist determined by the above-described processes effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier. In one embodiment, the mammalian MCH1 receptor agonist is not previously known.

This invention further provides a pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor antagonist determined by the above-described processes effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier. In one embodiment, the mammalian MCH1 receptor antagonist is not previously known.

This invention provides a method of screening a plurality of chemical compounds not known to activate a mammalian MCH1 receptor to identify a compound which activates the mammalian MCH1 receptor which comprises: (a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds not known to activate the mammalian MCH1 receptor, under conditions permitting activation of the mammalian MCH1 receptor; (b) determining whether the activity of the mammalian MCH1 receptor is increased in the presence of the compounds; and if so (c) separately determining whether the activation of the mammalian MCH1 receptor is increased by each compound included in the plurality of compounds, so as to thereby identify the compound which activates the mammalian MCH1 receptor. In one embodiment, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor.

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This invention provides a method of screening a plurality of chemical compounds not known to inhibit the activation of a mammalian MCH1 receptor to identify a compound which inhibits the activation of the mammalian MCH1 receptor, which comprises: (a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting activation of the mammalian MCH1 receptor; (b) determining whether the activation of the mammalian MCH1 receptor is reduced in the presence of the plurality of compounds, relative to the activation of the mammalian MCH1 receptor in the absence of the plurality of compounds; and if so (c) separately determining the inhibition of activation of the mammalian MCH1 receptor for each compound included in the plurality of compounds, so as to thereby identify the compound which inhibits the activation of the mammalian MCH1 receptor. In one embodiment, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor.

In one embodiment of the above-described methods, the cell is a mammalian cell. In another embodiment, the mammalian cell is non-neuronal in origin. In a further embodiment, the non-neuronal cell is a COS-7 cell, a 293 human embryonic kidney cell, a LM(tk-) cell or an NIH-3T3 cell.

This invention provides a pharmaceutical composition comprising a compound identified by the above-described methods effective to increase mammalian MCH1 receptor activity and a pharmaceutically acceptable carrier.

This invention also provides a pharmaceutical composition comprising a compound identified by the above-described methods effective to decrease mammalian MCH1 receptor activity and a pharmaceutically acceptable carrier.

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This invention further provides a method of measuring receptor activation in an oocyte expression system such as a *Xenopus* oocyte expression system or melanophore. In an embodiment, receptor activation is determined by measurement of ion channel activity. In another embodiment, receptor activation is measured by aequorin luminescence.

Expression of genes in *Xenopus* oocytes is well known in the art (Coleman, A., 1984; Masu, Y., et al., 1994) and is performed using microinjection of native mRNA or in vitro synthesized mRNA into frog oocytes. The preparation of in vitro synthesized mRNA can be performed by various standard techniques (Sambrook, et al. 1989) including using T7 polymerase with the mCAP RNA mapping kit (Stratagene).

This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor agonist effective to treat the abnormality. In separate embodiments, the abnormality is a regulation of a steroid or pituitary hormone disorder, an epinephrine release disorder, a gastrointestinal disorder, a cardiovascular disorder, an electrolyte balance disorder, hypertension, diabetes, a respiratory disorder, asthma, a reproductive function disorder, an immune disorder, an endocrine disorder, a musculoskeletal disorder, a neuroendocrine disorder, a cognitive disorder, a memory disorder such as Alzheimer's disease, a sensory modulation and transmission disorder, a motor coordination disorder, a sensory integration disorder, a motor integration disorder, a dopaminergic function disorder such as Parkinson's disease, a sensory transmission disorder, an olfaction disorder, a sympathetic innervation disorder, an affective disorder such as depression, a stress-related disorder, a fluid-balance disorder, a urinary disorder such as urinary incontinence, a seizure disorder, pain, psychotic behavior such as

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schizophrenia, morphine tolerance, opiate addiction or migraine.

This invention provides a method of treating an abnormality in a subject wherein the abnormality is alleviated by decreasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor antagonist effective to treat the abnormality. In separate embodiments, the abnormality is a regulation of a steroid or pituitary hormone disorder, an epinephrine release disorder, a gastrointestinal disorder, a cardiovascular disorder, an electrolyte balance disorder, hypertension, diabetes, a respiratory disorder, asthma, a reproductive function disorder, an immune disorder, an endocrine disorder, a musculoskeletal disorder, a neuroendocrine disorder, a cognitive disorder, a memory disorder such as Alzheimer's disease, a sensory modulation and transmission disorder, a motor coordination disorder, a sensory integration disorder, a motor integration disorder, a dopaminergic function disorder such as Parkinson's disease, a sensory transmission disorder, an olfaction disorder, a sympathetic innervation disorder, an affective disorder such as depression, a stress-related disorder, a fluid-balance disorder, a urinary disorder such as urinary incontinence, a seizure disorder, pain, psychotic behavior such as schizophrenia, morphine tolerance, opiate addiction or migraine.

This invention provides a process for making a composition of matter which specifically binds to a mammalian MCH1 receptor which comprises identifying a chemical compound using any of the processes described herein for identifying a compound which binds to and/or activates or inhibits activation of a mammalian MCH1 receptor and then synthesizing the chemical compound or a novel structural and functional analog or homolog thereof. In one embodiment, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH

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or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor.

This invention further provides a process for preparing a composition which comprises admixing a pharmaceutically acceptable carrier and a therapeutically effective amount of a chemical compound identified by any of the processes described herein for identifying a compound which binds to and/or activates or inhibits activation of a mammalian MCH1 receptor or a novel structural and functional analog or homolog thereof. In one embodiment, the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof. In another embodiment, the mammalian MCH1 receptor is a rat MCH1 receptor.

This invention provides a process for determining whether a chemical compound is a human MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH, and the cells do not express the MCH1 receptor prior to transfecting them.

This invention also provides a process for determining whether a chemical compound specifically binds to and inhibits activation of a human MCH1 receptor, which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1

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receptor, wherein such cells do not normally express the human MCH1 receptor and the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH. In one embodiment, the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

This invention further provides a method of screening a plurality of chemical compounds not known to inhibit the activation of a human MCH1 receptor to identify a compound which inhibits the activation of the human MCH1 receptor, which comprises:

- (a) contacting cells transfected with and expressing the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with the plurality of compounds in the presence of a known human MCH1 receptor agonist, under

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conditions permitting activation of the human MCH1 receptor, wherein the known MCH1 receptor agonist is MCH or a homolog or analog of MCH;

- 5       (b) determining whether the activation of the human MCH1 receptor is reduced in the presence of the plurality of compounds, relative to the activation of the human MCH1 receptor in the absence of the plurality of compounds; and if so
- 10      (c) separately determining the extent of inhibition of activation of the human MCH1 receptor for each compound included in the plurality of compounds, so as to thereby identify the compound which inhibits the activation of the human MCH1 receptor.
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In one embodiment of the above-described methods, the cell is an insect cell. In another embodiment, the cell is a mammalian cell. In still another embodiment, the cell is a mammalian cell which is nonneuronal in origin. In further embodiments, the cell is a COS-7 cell, a CHO cell, a 293 human embryonic kidney cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.

25      This invention provides a process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical

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compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor,  
5 wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a  
10 homolog or analog of MCH.

This invention further provides a process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting a membrane preparation from cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable  
15 for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second  
20 chemical compound is MCH or a homolog or analog of MCH.  
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This invention also provides a process for making a composition of matter which is a human MCH1 receptor

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antagonist which comprises identifying a chemical compound which is a human MCH1 receptor antagonist and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as a human MCH1 receptor antagonist by a process which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

This invention still further provides a process for making a composition of matter which specifically binds to and inhibits the activation of a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to and inhibits the activation of the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to and inhibiting the activation of the human MCH1 receptor by a process which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical

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compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH. In one embodiment, the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

This invention provides a process for preparing a composition which comprises identifying a chemical compound which specifically binds to a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or

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contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

5 This invention further provides a process for preparing a composition which comprises identifying a chemical compound which specifically binds to a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting a membrane preparation from cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

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30 This invention also provides a process for preparing a composition which comprises identifying a chemical compound which is a human MCH1 receptor antagonist, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as a human MCH1 receptor antagonist by a process which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation

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of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

This invention still further provides a process for preparing a composition which comprises identifying a chemical compound which specifically binds to and inhibits the activation of a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to and inhibiting activation of the human MCH1 receptor by a process which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or

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analog of MCH. In one embodiment, the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

In one embodiment of any of the above methods, the cell is an insect cell. In another embodiment, the cell is a mammalian cell. In another embodiment, the mammalian cell is nonneuronal in origin. In further embodiments, the nonneuronal cell is a COS-7 cell, a 293 human embryonic kidney cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.

Thus, once the gene for a targeted receptor subtype is cloned, it is placed into a recipient cell which then expresses the targeted receptor subtype on its surface. This cell, which expresses a single population of the targeted human receptor subtype, is then propagated resulting in the establishment of a cell line. This cell line, which constitutes a drug discovery system, is used in two different types of assays: binding assays and functional assays. In binding assays, the affinity of a compound for both the receptor subtype that is the target of a particular drug discovery program and other receptor subtypes that could be associated with side effects are measured. These measurements enable one to predict the potency of a compound, as well as the degree of selectivity that the compound has for the targeted receptor subtype over other receptor subtypes. The data obtained from binding assays also enable chemists to design compounds toward or away from one or more of the relevant subtypes, as appropriate, for optimal therapeutic efficacy. In functional assays, the nature of the response of the receptor subtype to the compound is determined. Data from the functional assays show whether the compound is acting to inhibit or enhance the activity of the receptor subtype, thus enabling

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pharmacologists to evaluate compounds rapidly at their ultimate human receptor subtypes targets permitting chemists to rationally design drugs that will be more effective and have fewer or substantially less severe side effects than 5 existing drugs.

Approaches to designing and synthesizing receptor subtype-selective compounds are well known and include traditional medicinal chemistry and the newer technology of 10 combinatorial chemistry, both of which are supported by computer-assisted molecular modeling. With such approaches, chemists and pharmacologists use their knowledge of the structures of the targeted receptor subtype and compounds determined to bind and/or activate or inhibit activation of 15 the receptor subtype to design and synthesize structures that will have activity at these receptor subtypes.

Combinatorial chemistry involves automated synthesis of a variety of novel compounds by assembling them using 20 different combinations of chemical building blocks. The use of combinatorial chemistry greatly accelerates the process of generating compounds. The resulting arrays of compounds are called libraries and are used to screen for compounds ("lead compounds") that demonstrate a sufficient level of 25 activity at receptors of interest. Using combinatorial chemistry it is possible to synthesize "focused" libraries of compounds anticipated to be highly biased toward the receptor target of interest.

Once lead compounds are identified, whether through the use 30 of combinatorial chemistry or traditional medicinal chemistry or otherwise, a variety of homologs and analogs are prepared to facilitate an understanding of the relationship between chemical structure and biological or 35 functional activity. These studies define structure activity relationships which are then used to design drugs with improved potency, selectivity and pharmacokinetic properties. Combinatorial chemistry is also used to rapidly

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generate a variety of structures for lead optimization. Traditional medicinal chemistry, which involves the synthesis of compounds one at a time, is also used for further refinement and to generate compounds not accessible by automated techniques. Once such drugs are defined the production is scaled up using standard chemical manufacturing methodologies utilized throughout the pharmaceutical and chemistry industry.

This invention will be better understood from the Experimental Details which follow. However, one skilled in the art will readily appreciate that the specific methods and results discussed are merely illustrative of the invention as described more fully in the claims which follow thereafter.

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EXPERIMENTAL DETAILS

Materials and methods

5       Cloning of human MCH1 receptor

Discovery of an Expressed Sequence Tag (EST) F07228 in GENEMBL Homologous to FB41a

A BLAST search of GENEMBL was performed with the GCG sequence analysis package (Genetics Computer Group, Madison, WI) using a Synaptic Pharmaceutical Corporation proprietary sequence, FB41a, as a query. This resulted in the identification of an EST (accession number F07228) with a high degree of homology to FB41a and somatostatin, opiate and galanin receptors.

15      Construction and Screening of a Human Hippocampal cDNA Library

Poly A+ RNA was purified from human hippocampal RNA (Clontech) using a FastTrack kit (Invitrogen, Corp.). DS-cDNA was synthesized from poly A+ RNA according to Gubler and Hoffman (1983) with minor modifications. The resulting cDNA was ligated to BstXI adaptors (Invitrogen, Corp.) and the excess adaptors removed by exclusion column chromatography. High molecular weight fractions of size-selected ds-cDNA were ligated in pEXJ.BS, an Okayama and Berg expression vector modified from pcEXV (Miller and Germain, 1986) to contain BstXI and other additional restriction sites. A total of  $2.2 \times 10^6$  independent clones with a mean insert size of 3.0 kb were generated. The library was plated on agar plates (ampicillin selection) and glycerol stocks for 450 pools of 5000 independent clones were prepared. Primary glycerol stocks were also grouped together in groups of approximately 10 to create superpools.

35      Cloning of the full-length sequence of MCH1  
Glycerol stocks of the superpools and primary pools from the

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human hippocampal cDNA library were screened by PCR with F07228 specific primers T579 and T580 using Taq DNA Polymerase (Boehringer-Mannheim, Indianapolis, IN) and the following PCR protocol: 94°C hold for 5 minutes; 40 cycles of 94°C for 2 minute, 68°C for 4 minutes; 7 minute hold at 68°C; 4°C hold until the samples are run on a gel. One positive primary pool 490, was successively divided into subpools, amplified in LB medium overnight and screened by PCR using primers T579 and T580. One positive subpool, 490-4-10-23 was plated on agar plates (ampicillin selection), and colonies were transferred to nitrocellulose membranes (Schleicher and Schuell, Keene, NH). Filters were hybridized for two days under high stringency conditions with  $10^6$  cpm/ml of a  $^{32}$ P-labeled cDNA probe, T581, designed against the F07228 EST sequence. Filters were washed and apposed to Biomax MS film (Kodak). Seven positive colonies were picked, streaked on LB-AMP plates, and grown overnight. Two individual colonies from each of the original seven were picked and subjected to vector-anchored PCR using the following primer pairs: T95, T580 and T94, T579. One positive colony, G1, was amplified overnight in TB and processed for plasmid purification. This plasmid was designated TL230 and sequenced on both strands with a Sequenase kit (US Biochemical, Cleveland, Ohio). Nucleotide and peptide sequence analysis were performed with GCG programs (Genetics Computer Group, Madison, WI). A HindIII-KpnI fragment of TL230 was subcloned into the mammalian expression vector pEXJ, and named TL231.

30 Primers and Probes:

TL579: 5'-GGGAACTCCACGGTCATCTTCGCGGT-3' (SEQ ID NO: 5)

TL580: 5'-TAGCGGTCAATGGCCATGGCGGTCAG-3' (SEQ ID NO: 6)

TL581:

35 5'-CTCCTGGGCATGCCCTTCATGATCCACCAGCTCATGGGCAATGGG-3'  
(SEQ ID NO: 7)

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TL94: 5'-CTTCTAGGCCTGTACGGAAGTGTAA-3' (SEQ ID NO: 8)

TL95: 5'-GTTGTGGTTGTCCAAACTCATCAATG-3' (SEQ ID NO: 9)

5       Isolation of a Fragment of a species homologue of TL231  
(human MCH1)

To obtain a fragment of a species homologue of TL231, the species genomic DNA (Clontech) may be amplified with a forward PCR primer corresponding to one of the TM regions of TL231 and a reverse primer corresponding to another TM region of TL231. PCR may be performed with the Expand Long Template PCR System (Boeringer Mannheim), for example, under the following conditions: 30 sec at 94°C, 1.5 min at 50°C, 1.5 min at 68 °C for 40 cycles, with a pre- and post-incubation of 5 min at 94°C and 7 min at 68°C, respectively. A band is isolated, subcloned using the TA cloning kit (Invitrogen), and sequenced. The sequence is run and analyzed on an ABI PRISM 377 BigDye Terminator Cycle Sequencing Kit Sequencer. Forward and reverse PCR primers are designed against this sequence and used to amplify a band from genomic DNA using, for example, the following conditions: 30 sec at 94°C, 1.5 min at 68°C for 35 cycles, with a pre- and post-incubation of 5 min at 94°C and 5 min at 68°C, respectively. The PCR product is subcloned using the TA cloning kit (Invitrogen). Miniprep cultures of transformants are prepared and sequenced as above.

30       Isolation of a full-length species homolog of TL231 (human  
MCH1)

A nucleic acid sequence encoding an MCH1 receptor may be isolated using standard molecular biology techniques and approaches such as those briefly described below:

35       Approach #1: To obtain a full-length MCH1 receptor, a cosmid library could be screened with a <sup>32</sup>P-labeled oligonucleotide probe.

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5       The full-length sequence may be obtained by sequencing this cosmid clone with additional sequencing primers. Since one intron is present in this gene the full-length intronless gene may be obtained from cDNA using standard molecular biology techniques. For example, a forward PCR primer designed in the 5'UT and a reverse PCR primer designed in the 3'UT may be used to amplify a full-length, intronless gene from cDNA. Standard molecular biology techniques could be used to subclone this gene into a mammalian expression  
10      vector.

15      Approach #2: Standard molecular biology techniques could be used to screen commercial cDNA phage libraries by hybridization under high stringency with a <sup>32</sup>P-labeled oligonucleotide probe. One may isolate a full-length MCH1 receptor by obtaining a plaque purified clone from the lambda libraries and then subjecting the clone to direct DNA sequencing. Alternatively, standard molecular biology techniques could be used to screen in-house cDNA plasmid  
20      libraries by PCR amplification of library pools using primers to the MCH1 sequence. A full-length clone could be isolated by Southern hybridization of colony lifts of positive pools with a <sup>32</sup>P-labeled oligonucleotide probe.

25      Approach #3: As yet another alternative method, one could utilize 3' and 5' RACE to generate PCR products from cDNA expressing MCH1 which contain the additional sequences of MCH1. These RACE PCR products could then be sequenced to determine the missing sequence. This new sequence could then be used to design a forward PCR primer in the 5'UT and a reverse primer in the 3'UT. These primers could then be used to amplify a full-length MCH1 clone from cDNA.  
30

Construction of Human MCH1 Mutants

35      The plasmid TL231 encodes three in frame methionine residues, any of which could potentially initiate translation of the MCH1 receptor. The ability of these

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residues to function in a heterologous expression system was examined by constructing mutants of TL231 in which one or more of the downstream methionine residues was mutated to alanine. Mutagenesis was performed using the QuickChange site-directed mutagenesis kit (Stratagene). Each 50 ul PCR reaction contained 10 mM KCl, 10 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 20 mM Tris-HCl (pH 8.8), 2 mM MgSO<sub>4</sub>, 0.1% Triton X-100, 0.1mg/ml nuclease-free BSA, 114 ng each of two mutagenesis primers (see below), 50 ng of plasmid DNA template (see below), 2.5 units of PfuTurbo DNA polymerase, and 1 ul of the proprietary dNTP mix provided in the kit. Thermocycling was performed with an Applied Biosystems 9700 machine using the following cycling parameters: one cycle of 95° for 30 seconds; eighteen cycles of 95° for 30 seconds, 55° for 1 minute, 68° for 2.5 minutes; a final hold at 4°. Next, 1 ul (10 units) of DpnI restriction enzyme was added to the mutagenesis reaction followed by incubation at 37° for 1 hour. A 2 ul aliquot of this digestion was used to transform 50 ul of E.coli XL1-Blue cells provided with the mutagenesis kit. Transformants were selected by their ability to grow at 37° on LB plates containing 100 ug/ml ampicillin. Single colonies which resulted from the overnight incubation of the plates were used to inoculate 2 ml cultures of LB-ampicillin and allowed to grow overnight at 37° with shaking. Miniprep DNA was prepared from these cultures using the Qiagen miniprep system and subjected to automated sequence analysis. This allowed both the confirmation of the desired mutation and the integrity of the remainder of the MCH1 coding sequence. After identification of a correctly mutated clone, a large scale DNA prep was prepared using a Qiagen megaprep column.

To create the clone encoding only the M70A mutation, the template DNA was TL231 and the mutagenesis primers were RP192 and RP193. This clone is designated R106 (SEQ ID NO:

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16) and encodes only the first two potential start codons (See Figure 12). To create the clone encoding both the M6A and the M70A mutations, the template DNA was R106 and the mutagenesis primers were RP190 and RP191. The resulting 5 clone is designated R114 (SEQ ID NO: 17) and encodes only first start codon (See Figure 12).

If desired, the same mutagenesis technology can be employed 10 to construct additional MCH1 mutants that encode other combinations of the available methionine residues. The mutation M1A could be constructed using primers X1 and X2. Such a change would eliminate the first methionine but retain the two downstream residues. Likewise, the double 15 mutation M1A, M70A could be constructed by sequentially using primer pairs X1/ X2 and RP192/RP193. This would create a gene in which only the second methionine was left intact.

Primers used in the generation of hMCH1 mutant receptor constructs:

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Mutant Primer Primer Sequence

R106 RP192 5' CGGCAC TGGCTGGCGGACCTGGAAGCCCTCG 3' (SEQ  
ID NO: 18)

M70A) RP193 5' CGAGGGCTTCCAGGTCCGCCAGCCAGTGCCG 3'  
(SEQ ID NO: 19)

25

R114 RP190 5' ATGTCAGTGGGAGCCCGCAAGAAGGGAGTGGG 3' (SEQ  
ID NO: 20)

(M6A, RP191 5' CCCACTCCCTTCTCGCGGCTCCC ACTGACAT 3'  
30 M70A) (SEQ ID NO: 21)

(M1A) X1 5' TAATGTGTCTAGGTGGCGTCAGTGGGAGCCATG 3' (SEQ  
ID NO: 22)  
X2 5' CATGGCTCCACTGACGCCACCTAGACACATTA 3' (SEQ  
35 ID NO: 23)

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Construction of a short form of the human MCH1 receptor

A short form of the human MCH1 receptor expressing only the  
5 most downstream of the three potential initiating  
methionines was generated as follows. TL231 was amplified  
with BB1122 (a forward primer beginning 10 nucleotides  
upstream of the third methionine in TL231, and also  
incorporating a *Hind*III site) and BB1123 (a reverse primer  
10 in the second transmembrane domain) and the resulting  
product digested with *Hind*III and *Bgl*IIA. PCR was performed  
with the Expand Long Template PCR System (Roche Molecular  
Biochemicals, Indianapolis, IN) under the following  
conditions: 20 seconds at 94°C, 1 minute at 68°C for 40  
15 cycles, with a pre- and post-incubation of 5 minutes at 94°C  
and 7 minutes at 68°C respectively. The 270 bp product was  
gel purified and ligated to a 4 kb *Hind*III/*Bgl*II  
restriction fragment from TL231. The resulting construct  
was named BO120 (SEQ ID NO: 24).

20 Primers used in the construction of the truncated human MCH1  
receptor:

BB1122 5'- TGACACTAACGCTTCACTGGCTGGATGGACCTGGAAGC -3' (SEQ  
25 ID NO: 25)

BB1123 5'- GCCCAGGAGAAAGAGGGAGATCTAC -3' (SEQ ID NO: 26)

30 Host cells

A broad variety of host cells can be used to study  
heterologously expressed proteins. These cells include but  
are not restricted to assorted mammalian lines such as; Cos-  
7, CHO, LM(tk-), HEK293, etc.; insect cell lines such as;  
35 Sf9, Sf21, etc.; amphibian cells such as *xenopus* oocytes;  
and others.

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5 COS-7 cells are grown on 150 mm plates in DMEM with supplements (Dulbecco's Modified Eagle Medium with 10% bovine calf serum, 4 mM glutamine, 100 units/ml penicillin/100 µg/ml streptomycin) at 37°C, 5% CO<sub>2</sub>. Stock plates of COS-7 cells are trypsinized and split 1:6 every 3-4 days.

10 Human embryonic kidney 293 cells are grown on 150 mm plates in DMEM with supplements (10% bovine calf serum, 4 mM glutamine, 100 units/ml penicillin/100 µg/ml streptomycin) at 37°C, 5% CO<sub>2</sub>. Stock plates of 293 cells are trypsinized and split 1:6 every 3-4 days.

15 Mouse fibroblast LM(tk-) cells are grown on 150 mm plates in D-MEM with supplements (Dulbecco's Modified Eagle Medium with 10% bovine calf serum, 4 mM glutamine, 100 units/ml penicillin/100 µg/ml streptomycin) at 37°C, 5% CO<sub>2</sub>. Stock plates of LM(tk-) cells are trypsinized and split 1:10 every 20 3-4 days.

25 Chinese hamster ovary (CHO) cells were grown on 150 mm plates in HAM's F-12 medium with supplements (10% bovine calf serum, 4 mM L-glutamine and 100 units/ml penicillin/100 µg/ml streptomycin) at 37°C, 5% CO<sub>2</sub>. Stock plates of CHO cells are trypsinized and split 1:8 every 3-4 days.

30 Mouse embryonic fibroblast NIH-3T3 cells are grown on 150 mm plates in Dulbecco's Modified Eagle Medium (DMEM) with supplements (10% bovine calf serum, 4 mM glutamine, 100 units/ml penicillin/100 µg/ml streptomycin) at 37°C, 5% CO<sub>2</sub>. Stock plates of NIH-3T3 cells are trypsinized and split 1:15 every 3-4 days.

35 Sf9 and Sf21 cells are grown in monolayers on 150 mm tissue culture dishes in TMN-FH media supplemented with 10% fetal calf serum, at 27°C, no CO<sub>2</sub>. High Five insect cells are grown on 150 mm tissue culture dishes in Ex-Cell 400™ medium

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supplemented with L-Glutamine, also at 27°C, no CO<sub>2</sub>.

In some cases, cell lines that grow as adherent monolayers can be converted to suspension culture to increase cell yield and provide large batches of uniform assay material for routine receptor screening projects.

Xenopus oocytes can also be used as a host system for transient expression of heterologous proteins. Their maintenance and usage is described in the electrophysiological methods section that follows.

Transient expression

DNA encoding proteins to be studied can be transiently expressed in a variety of mammalian, insect, amphibian and other cell lines by several methods including but not restricted to; calcium phosphate-mediated, DEAE-dextran mediated, Liposomal-mediated, viral-mediated, electroporation-mediated and microinjection delivery. Each of these methods may require optimization of assorted experimental parameters depending on the DNA, cell line, and the type of assay to be subsequently employed.

A typical protocol for the calcium phosphate method as applied to LM(tk-) cells is described as follows; Adherent cells are harvested approximately twenty-four hours before transfection and replated at a density of 1-2 x 10<sup>5</sup> cells/cm<sup>2</sup> in a 100 mm tissue culture dish and allowed to incubate over night at 37°C at 5% CO<sub>2</sub>. 250 µl of a mixture of CaCl<sub>2</sub> and DNA (20 µg DNA in 250 mM CaCl<sub>2</sub>) is added to a 5 ml plastic tube and 250 ul of 2X HBS (250 mM NaCl, 10 mM KCl, 1.5 mM Na<sub>2</sub>HPO<sub>4</sub>, 12 mM dextrose, 50 mM HEPES) is slowly added with gentle mixing. The mixture is allowed to incubate for 20 minutes at room temperature to allow a DNA precipitate to form. The cells are then washed with complete medium, 10 ml of culture medium is added to each plate, followed by addition of the DNA precipitate. The cells are then

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incubated for 24 to 48 hours at 37°C at 5% CO<sub>2</sub>.

A typical protocol for the DEAE-dextran method as applied to Cos-7 cells is described as follows; Cells to be used for transfection are split 24 hours prior to the transfection to provide flasks which are 70-80% confluent at the time of transfection. Briefly, 8 µg of receptor DNA plus 8 µg of any additional DNA needed (e.g. G<sub>a</sub> protein expression vector, reporter construct, antibiotic resistance marker, mock vector, etc.) are added to 9 ml of complete DMEM plus DEAE-dextran mixture (10 mg/ml in PBS). Cos-7 cells plated into a T225 flask (sub-confluent) are washed once with PBS and the DNA mixture is added to each flask. The cells are allowed to incubate for 30 minutes at 37°C, 5% CO<sub>2</sub>. Following the incubation, 36 ml of complete DMEM with 80 µM chloroquine is added to each flask and allowed to incubate an additional 3 hours. The medium is then aspirated and 24 ml of complete medium containing 10% DMSO for exactly 2 minutes and then aspirated. The cells are then washed 2 times with PBS and 30 ml of complete DMEM added to each flask. The cells are then allowed to incubate over night. The next day the cells are harvested by trypsinization and reseeded as needed depending upon the type of assay to be performed.

A typical protocol for liposomal-mediated transfection as applied to CHO cells is described as follows; Cells to be used for transfection are split 24 hours prior to the transfection to provide flasks which are 70-80% confluent at the time of transfection. A total of 10µg of DNA which may include varying ratios of receptor DNA plus any additional DNA needed (e.g. G<sub>a</sub> protein expression vector, reporter construct, antibiotic resistance marker, mock vector, etc.) is used to transfet each 75 cm<sup>2</sup> flask of cells. Liposomal mediated transfection is carried out according to the manufacturer's recommendations (LipofectAMINE, GibcoBRL, Bethesda, MD). Transfected cells are harvested 24 h post transfection and used or reseeded according the requirements

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of the assay to be employed.

A typical protocol for the electroporation method as applied to Cos-7 cells is described as follows; Cells to be used for transfection are split 24 hours prior to the transfection to provide flasks which are subconfluent at the time of transfection. The cells are harvested by trypsinization resuspended in their growth media and counted.  $4 \times 10^4$  cells are suspended in 300  $\mu\text{l}$  of DMEM and placed into an electroporation cuvette. 8  $\mu\text{g}$  of receptor DNA plus 8  $\mu\text{g}$  of any additional DNA needed (e.g. G<sub>a</sub> protein expression vector, reporter construct, antibiotic resistance marker, mock vector, etc.) is added to the cell suspension, the cuvette is placed into a BioRad Gene Pulser and subjected to an electrical pulse (Gene Pulser settings: 0.25 kV voltage, 950  $\mu\text{F}$  capacitance). Following the pulse, 800  $\mu\text{l}$  of complete DMEM is added to each cuvette and the suspension transferred to a sterile tube. Complete medium is added to each tube to bring the final cell concentration to  $1 \times 10^5$  cells/100  $\mu\text{l}$ . The cells are then plated as needed depending upon the type of assay to be performed.

A typical protocol for viral mediated expression of heterologous proteins is described as follows for baculovirus infection of insect Sf9 cells. The coding region of DNA encoding the receptor disclosed herein may be subcloned into pBlueBacIII into existing restriction sites or sites engineered into sequences 5' and 3' to the coding region of the polypeptides. To generate baculovirus, 0.5  $\mu\text{g}$  of viral DNA (BaculoGold) and 3  $\mu\text{g}$  of DNA construct encoding a polypeptide may be co-transfected into  $2 \times 10^6$  *Spodoptera frugiperda* insect Sf9 cells by the calcium phosphate coprecipitation method, as outlined in by Pharmingen (in "Baculovirus Expression Vector System: Procedures and Methods Manual"). The cells then are incubated for 5 days at 27°C. The supernatant of the co-transfection plate may be collected by centrifugation and the recombinant virus plaque purified. The procedure to infect cells with virus, to

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5 prepare stocks of virus and to titer the virus stocks are as described in Pharmingen's manual. Similar principals would in general apply to mammalian cell expression via retro-viruses, Simliki forest virus and double stranded DNA viruses such as adeno-, herpes-, and vaccinia-viruses, and the like.

10 Microinjection of cRNA encoding for proteins of interest is useful for the study of protein function in xenopus oocytes as well as cultured mammalian cells. A typical protocol for the preparation of cRNA and injection into xenopus oocytes can be found in the following electrophysiology section.

Stable expression

15 Heterologous DNA can be stably incorporated into host cells, causing the cell to perpetually express a foreign protein. Methods for the delivery of the DNA into the cell are similar to those described above for transient expression but require the co-transfection of an ancillary gene to confer drug resistance on the targeted host cell. The ensuing drug resistance can be exploited to select and maintain cells that have taken up the heterologous DNA. An assortment of resistance genes are available including but not restricted to Neomycin, Kanamycin, and Hygromycin. For 20 the purposes of receptor studies, stable expression of a heterologous receptor protein is carried out in, but not necessarily restricted to, mammalian cells including, CHO, HEK293, LM(tk-), etc.

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30 Cell membrane preparation

For binding assays, pellets of transfected cells are suspended in ice-cold buffer (20 mM Tris.HCl, 5 mM EDTA, pH 7.4) and homogenized by sonication for 7 sec. The cell lysates are centrifuged at 200 x g for 5 min at 4°C. The supernatants are then centrifuged at 40,000 x g for 20 min at 4°C. The resulting pellets are washed once in the homogenization buffer and suspended in binding buffer (see methods for radioligand binding). Protein concentrations are

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determined by the method of Bradford (1976) using bovine serum albumin as the standard. Binding assays are usually performed immediately, however it is possible to prepare membranes in batch and store frozen in liquid nitrogen for future use.

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Radioligand binding assays

Cells may be screened for the presence of endogenous human receptor by radioligand binding (described in detail below).  
10 Cells with either no or a low level of the endogenous human receptor disclosed herein may be transfected with the exogenous receptor.

MCH1 binding experiments with membranes (20-40 µg membrane protein) from transfected cells are performed with 0.1 nM [<sup>125</sup>I]Phe<sup>13</sup>-Tyr<sup>19</sup>-MCH (Custom labeled by NEN) using incubation buffer consisting of 50mM Tris pH 7.4, 10mM MgCl<sub>2</sub>, 2 µg/ml aprotinin, 0.5mM PMSF and 50 µg/ml bacitracin. Binding is performed at 25°C for 1 hr. Incubations are terminated by rapid vacuum filtration over GF/C glass fiber filters, presoaked in 5% PEI using 50 mM Tris pH 7.4 containing 0.01% triton X-100 as wash buffer. In all experiments nonspecific binding is defined using 10 µM unlabeled MCH.

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Functional assays

Cells may be screened for the presence of endogenous mammalian receptor using functional assays (described in detail below). Cells with no or a low level of endogenous receptor present may be transfected with the exogenous receptor for use in the following functional assays.

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A wide spectrum of assays can be employed to screen for receptor activation. These range from traditional measurements of phosphatidyl inositol, cAMP, Ca<sup>++</sup>, and K<sup>+</sup>, for example; to systems measuring these same second messengers but which have been modified or adapted to be higher throughput, more generic, and more sensitive; to cell based platforms reporting more general cellular events

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resulting from receptor activation such as metabolic changes, differentiation, and cell division/proliferation, for example; to high level organism assays which monitor complex physiological or behavioral changes thought to be involved with receptor activation including cardiovascular, analgesic, orexigenic, anxiolytic, and sedation effects, for example.

Cyclic AMP (cAMP) assay  
The receptor-mediated stimulation or inhibition of cyclic AMP (cAMP) formation may be assayed in cells expressing the mammalian receptors. Cells are plated in 96-well plates and incubated in Dulbecco's phosphate buffered saline (PBS) supplemented with 10 mM HEPES, 1mM isobutylmethylxanthine for 20 min at 37°C, in 5% CO<sub>2</sub>. Test compounds are added with or without 10 μM forskolin and incubated for an additional 10 min at 37°C. The medium is then aspirated and the reaction stopped by the addition of 100 mM HCl. The plates are stored at 4°C for 15 min, and the cAMP content in the stopping solution measured by radioimmunoassay. Radioactivity may be quantified using a gamma counter equipped with data reduction software.

Arachidonic acid release assay  
Cells expressing the mammalian receptor are seeded into 96 well plates and grown for 3 days in HAM's F-12 with supplements. [<sup>3</sup>H]-arachidonic acid (specific activity = 0.75 μCi/ml) is delivered as a 100 μL aliquot to each well and samples were incubated at 37° C, 5% CO<sub>2</sub> for 18 hours. The labeled cells are washed three times with 200 μL HAM's F-12. The wells are then filled with medium (200 μL) and the assay is initiated with the addition of peptides or buffer (22 μL). Cells are incubated for 30 min at 37°C, 5% CO<sub>2</sub>. Supernatants are transferred to a microtiter plate and evaporated to dryness at 75°C in a vacuum oven. Samples are then dissolved and resuspended in 25 μL distilled water. Scintillant (300 μL) is added to each well and samples are counted for <sup>3</sup>H in a Trilux plate reader. Data are analyzed

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using nonlinear regression and statistical techniques available in the GraphPAD Prism package (San Diego, CA).

Intracellular calcium mobilization assay

5       The intracellular free calcium concentration may be measured by microspectrofluorometry using the fluorescent indicator dye Fura-2/AM (Bush et al, 1991). Cells are seeded onto a 35 mm culture dish containing a glass coverslip insert, washed with HBS and loaded with 100  $\mu$ L of Fura-2/AM (10  $\mu$ M) for 20 to 40 min. After washing with HBS to remove the Fura-2/AM solution, cells are equilibrated in HBS for 10 to 10 min. Cells are then visualized under the 40X objective of a Leitz Fluovert FS microscope and fluorescence emission is determined at 510 nM with excitation wavelengths alternating between 340 nM and 380 nM. Raw fluorescence data are converted to calcium concentrations using standard calcium concentration curves and software analysis techniques.

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20       Inositol phosphate assay

Guidelines for cell preparation and assay of the second messenger inositol phosphate (IP) are described below for a typical protocol involving transiently transfected Cos-7 cells; For a 96 well microplate format assay, cells are plated at 70,000 cells per well and allowed to incubate for 24 hours after the transfection procedure. The cells are then labeled with 0.5  $\mu$ Ci [ $^3$ H]myo-inositol per micro-well over night at 37°C, 5% CO<sub>2</sub>. Immediately before the assay, the medium is removed and replaced with 90  $\mu$ l PBS containing 10 mM LiCl. The plates are then incubated for 15 minutes at 37°C, 5% CO<sub>2</sub>. Following the incubation, the transfectants are challenged with agonist (10  $\mu$ l/well; 10X concentration) for 30 minutes at 37°C, 5% CO<sub>2</sub>. The challenge is terminated and the cells lysed by the addition of 100  $\mu$ l cold 5% v/v trichloroacetic acid (TCA), followed by an incubation at 4°C for greater than 30 minutes. Total IPs are isolated from the lysate by ion exchange chromatography. Briefly, the

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lys<sup>5</sup>ed contents of the wells are transferred to a Multiscreen HV filter plate (Millipore) containing 100  $\mu$ l Dowex AG1-X8 suspension (50% v/v, water:resin) (200-400 mesh, formate form). The filter plates are placed on a vacuum manifold to wash and elute the resin bed. Each well is first washed 2 times with 200  $\mu$ l 5 mM myoinositol. Total [<sup>3</sup>H]IPs are eluted with 75  $\mu$ l of 1.2 M ammonium formate/0.1 M formic acid into Wallac 96-well plates. 200  $\mu$ l of SuperMix scintillation cocktail is added to each well, mixed well, allowed to equilibrate and counted on a Micro Beta Trilux scintillation counter. (Note: The assay may be scaled to a 24 well format by simple adjustment of reagent volumes and employing individual chromatographic columns.)

15 GTP $\gamma$ S functional assay

Membranes from cells transfected with the mammalian receptors are suspended in assay buffer (50 mM Tris, 100 mM NaCl, 5 mM MgCl<sub>2</sub>, pH 7.4) supplemented with 0.2% BSA and 10  $\mu$ M GDP. Membranes are incubated on ice for 20 minutes, transferred to a 96-well Millipore microtiter GF/C filter plate and mixed with GTP $\gamma$ <sup>35</sup>S (e.g., 250,000 cpm/sample, specific activity ~1000 Ci/mmol) plus or minus GTP $\gamma$ S (final concentration = 100  $\mu$ M). Final membrane protein concentration = 90  $\mu$ g/ml. Samples are incubated in the presence or absence of MCH (final concentration = 1  $\mu$ M) for 30 min. at room temperature, then filtered on a Millipore vacuum manifold and washed three times with cold assay buffer. Samples collected in the filter plate are treated with scintillant and counted for <sup>35</sup>S in a Trilux (Wallac) liquid scintillation counter. It is expected that optimal results are obtained when the mammalian receptor membrane preparation is derived from an appropriately engineered heterologous expression system, i.e., an expression system resulting in high levels of expression of the mammalian receptor and/or expressing G-proteins having high turnover rates (for the exchange of GDP for GTP). GTP $\gamma$ S assays are well-known in the art, and it is expected that variations on

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the method described above, such as are described by e.g., Tian et al. (1994) or Lazareno and Birdsall (1993), may be used by one of ordinary skill in the art.

5       Transcription assay

Guidelines for cell preparation and assay of receptor mediated transcription of Cos-7 cells transiently transfected by the DEAE-dextran method in a 96 microwell format is as follows; The c-fos- $\beta$ -gal promoter/reporter construct used for these studies consists of the cfos promoter region (-384 to +19) (Schilling et al 1991, Yalkinoglu et al, 1995) inserted upstream of  $\beta$ -galactosidase cDNA containing expression vector pNASS $\beta$  (Clontech). Transcription activity is measured by assay of  $\beta$ -galactosidase enzyme activity as detected in a colorimetric assay. Forty-eight hours following transient transfection, the medium is removed and replaced with medium containing drug (e.g. MCH) typically at a concentration of 10  $\mu$ M. The cells are allowed to incubate at 37°C, 5% CO<sub>2</sub> for at least 18 hours, after which the medium is aspirated and the cells washed with 200  $\mu$ l PBS/well. The cells are then lysed with 100  $\mu$ l AB buffer (100 mM Sodium Phosphate buffer, pH 8.0, 2 mM MgSO<sub>4</sub>, 0.1 mM MnCl<sub>2</sub>) for 10 minutes at room temperature. 100  $\mu$ l of AB/Tx/ $\beta$ -mercaptoethanol (AB buffer with 0.5% Triton X-100, 40 mM  $\beta$ -mercaptoethanol) is then added to each well and the lysate allowed to incubate an additional 10 minutes at room temperature. The enzymatic color reaction is initiated by the addition of the substrate, ONPG/AB (4 mg/ml O-nitrophenyl- $\beta$ -D-galactopyranoside in AB buffer). The reaction is allowed to proceed for 30 minutes or until yellow color becomes evident. Measurement of optical density is taken at 405 nm using a Dynatech microplate reader.

35      MAP kinase assay

MAP kinase (mitogen activated kinase) may be monitored to evaluate receptor activation. MAP kinase is activated by multiple pathways in the cell. A primary mode of activation

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5 involves the ras/raf/MEK/MAP kinase pathway. Growth factor (tyrosine kinase) receptors feed into this pathway via SHC/Grb-2/SOS/ras. Gi coupled receptors are also known to activate ras and subsequently produce an activation of MAP kinase. Receptors that activate phospholipase C (Gq and G11) produce diacylglycerol (DAG) as a consequence of phosphatidyl inositol hydrolysis. DAG activates protein kinase C which in turn phosphorylates MAP kinase.

10 MAP kinase activation can be detected by several approaches. One approach is based on an evaluation of the phosphorylation state, either unphosphorylated (inactive) or phosphorylated (active). The phosphorylated protein has a slower mobility in SDS-PAGE and can therefore be compared with the unstimulated protein using Western blotting.

15 Alternatively, antibodies specific for the phosphorylated protein are available (New England Biolabs) which can be used to detect an increase in the phosphorylated kinase. In either method, cells are stimulated with the mitogen and then extracted with Laemmli buffer. The soluble fraction is applied to an SDS-PAGE gel and proteins are transferred electrophoretically to nitrocellulose or Immobilon. Immunoreactive bands are detected by standard Western blotting technique. Visible or chemiluminescent signals are recorded on film and may be quantified by densitometry.

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Another approach is based on evaluation of the MAP kinase activity via a phosphorylation assay. Cells are stimulated with the mitogen and a soluble extract is prepared. The extract is incubated at 30°C for 10 min with gamma-32-ATP, an ATP regenerating system, and a specific substrate for MAP kinase such as phosphorylated heat and acid stable protein regulated by insulin, or PHAS-I. The reaction is terminated by the addition of H<sub>3</sub>PO<sub>4</sub> and samples are transferred to ice. An aliquot is spotted onto Whatman P81 chromatography paper, which retains the phosphorylated protein. The chromatography paper is washed and counted for <sup>32</sup>P in a liquid scintillation counter. Alternatively, the cell

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extract is incubated with gamma-<sup>32</sup>-ATP, an ATP regenerating system, and biotinylated myelin basic protein bound by streptavidin to a filter support. The myelin basic protein is a substrate for activated MAP kinase. The phosphorylation reaction is carried out for 10 min at 30°C. The extract can then be aspirated through the filter, which retains the phosphorylated myelin basic protein. The filter is washed and counted for <sup>32</sup>P by liquid scintillation counting.

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Cell proliferation assay

Activation of a G protein coupled receptor may lead to a mitogenic or proliferative response which can be monitored via [<sup>3</sup>H]-thymidine uptake. When cultured cells are incubated with [<sup>3</sup>H]-thymidine, the thymidine translocates into the nuclei where it is phosphorylated to thymidine triphosphate. The nucleotide triphosphate is then incorporated into the cellular DNA at a rate that is proportional to the rate of cell growth. Typically, cells are grown in culture for 1-3 days. Cells are forced into quiescence by the removal of serum for 24 hrs. A mitogenic agent is then added to the media. 24 hrs later, the cells are incubated with [<sup>3</sup>H]-thymidine at specific activities ranging from 1 to 10 µCi/ml for 2-6 hrs. Harvesting procedures may involve trypsinization and trapping of cells by filtration over GF/C filters with or without a prior incubation in TCA to extract soluble thymidine. The filters are processed with scintillant and counted for <sup>3</sup>H by liquid scintillation counting. Alternatively, adherent cells are fixed in MeOH or TCA, washed in water, and solubilized in 0.05% deoxycholate/0.1 N NaOH. The soluble extract is transferred to scintillation vials and counted for <sup>3</sup>H by liquid scintillation counting.

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Methods for recording currents in Xenopus oocytes

Female *Xenopus laevis* (*Xenopus-1*, Ann Arbor, MI) are anesthetized in 0.2% tricain (3-aminobenzoic acid ethyl

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ester, Sigma Chemical Corp.) and a portion of ovary is removed using aseptic technique (Quick and Lester, 1994). Oocytes are defolliculated using 2 mg/ml collagenase (Worthington Biochemical Corp., Freehold, NJ) in a solution containing 87.5 mM NaCl, 2 mM KCl, 2 mM MgCl<sub>2</sub> and 5 mM HEPES, pH 7.5. Oocytes may be injected (Nanoject, Drummond Scientific, Broomall, PA) with mammalian mRNA. Other oocytes may be injected with a mixture of mammalian mRNA and mRNA encoding the genes for G-protein-activated inward rectifiers (GIRK1 and GIRK4, U.S. Patent Nos. 5,734,021 and 5,728,535). Genes encoding G-protein inwardly rectifying K<sup>+</sup> (GIRK) channels 1 and 4 (GIRK1 and GIRK4) were obtained by PCR using the published sequences (Kubo et al., 1993; Dascal et al., 1993; Krapivinsky et al., 1995 and 1995b) to derive appropriate 5' and 3' primers. Human heart cDNA was used as template together with the primers  
5'-CGCGGATCCATTATGTCTGCACCCGAAGGAAATTG-3' (SEQ ID NO: 10)  
and  
5'-CGCGAATTCTTATGTGAAGCGATCAGAGTTCATTTTC-3' (SEQ ID NO: 11)  
for GIRK1 and  
5'-GCGGGATCCGCTATGGCTGGTGATTCTAGGAATG-3' (SEQ ID NO: 12) and  
5'- CCGGAATTCCCCTCACACCGAGGCCCTGG-3' (SEQ ID NO: 13) for  
GIRK4.  
In each primer pair, the upstream primer contained a BamHI site and the downstream primer contained an EcoRI site to facilitate cloning of the PCR product into pcDNA1-Amp (Invitrogen). The transcription template for the mammalian receptor may be similarly obtained. mRNAs are prepared from separate DNA plasmids containing the complete coding regions of the mammalian receptor, GIRK1, and GIRK4. Plasmids are linearized and transcribed using the T7 polymerase ("Message Machine", Ambion). Alternatively, mRNA may be translated from a template generated by PCR, incorporating a T7 promoter and a poly A' tail. Each oocyte receives 2 ng each of GIRK1 and GIRK4 mRNA in combination with 25 ng of mammalian receptor mRNA. After injection of mRNA, oocytes are incubated at 16° C on a rotating platform for 3-8 days. Dual electrode voltage clamp ("GeneClamp", Axon Instruments

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Inc., Foster City, CA) is performed using 3 M KCl-filled glass microelectrodes having resistances of 1-3 Mohms. Unless otherwise specified, oocytes are voltage clamped at a holding potential of -80 mV. During recordings, oocytes are bathed in continuously flowing (2-5 ml/min) medium containing 96 mM NaCl, 2 mM KCl, 2 mM CaCl<sub>2</sub>, 2 mM MgCl<sub>2</sub>, and 5 mM HEPES, pH 7.5 ("ND96"), or, in the case of oocytes expressing GIRK1 and GIRK4, elevated K<sup>+</sup> containing 96 mM KCl, 2 mM NaCl, 2 mM CaCl<sub>2</sub>, 2 mM MgCl<sub>2</sub>, and 5 mM HEPES, pH 7.5 ("hK"). Drugs are applied by switching from a series of gravity fed perfusion lines.

Heterologous expression of GPCRs in *Xenopus* oocytes has been widely used to determine the identity of signaling pathways activated by agonist stimulation (Gundersen et al., 1983; Takahashi et al., 1987). Activation of the phospholipase C (PLC) pathway is assayed by applying test compound in ND96 solution to oocytes previously injected with mRNA for the mammalian receptor and observing inward currents at a holding potential of -80 mV. The appearance of currents that reverse at -25 mV and display other properties of the Ca<sup>++</sup>-activated Cl<sup>-</sup> (chloride) channel is indicative of mammalian receptor-activation of PLC and release of IP<sub>3</sub> and intracellular Ca<sup>++</sup>. Such activity is exhibited by GPCRs that couple to G<sub>q</sub>.

Measurement of inwardly rectifying K<sup>+</sup> (potassium) channel (GIRK) activity is monitored in oocytes that have been co-injected with mRNAs encoding the mammalian receptor, GIRK1, and GIRK4. The two GIRK gene products co-assemble to form a G-protein activated potassium channel known to be activated (i.e., stimulated) by a number of GPCRs that couple to G<sub>i</sub> or G<sub>o</sub> (Kubo et al., 1993; Dascal et al., 1993). Oocytes expressing the mammalian receptor plus the two GIRK subunits are tested for test compound responsivity by measuring K<sup>+</sup> currents in elevated K<sup>+</sup> solution (hK). Activation of inwardly rectifying currents that are sensitive to 300 μM Ba<sup>++</sup> signifies the mammalian receptor

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coupling to a G<sub>i</sub> or G<sub>e</sub> pathway in the oocytes.

Receptor/G protein co-transfection studies

A strategy for determining whether MCH1 can couple preferentially to selected G proteins involves co-transfection of MCH1 receptor cDNA into a host cell together with the cDNA for a G protein alpha sub-unit. Examples of G alpha sub-units include members of the G<sub>ai</sub>/G<sub>ao</sub> class (including G<sub>at2</sub> and G<sub>az</sub>), the G<sub>aq</sub> class, the G<sub>as</sub> class, and the G<sub>a12/13</sub> class. A typical procedure involves transient transfection into a host cell such as COS-7. Other host cells may be used. A key consideration is whether the cell has a downstream effector (a particular adenylyl cyclase, phospholipase C, or channel isoform, for example) to support a functional response through the G protein under investigation. G protein beta gamma sub-units native to the cell are presumed to complete the G protein heterotrimer; otherwise specific beta and gamma sub-units may be co-transfected as well. Additionally, any individual or combination of alpha, beta, or gamma subunits may be co-transfected to optimize the functional signal mediated by the receptor.

The receptor/G alpha co-transfected cells are evaluated in a binding assay, in which case the radioligand binding may be enhanced by the presence of the optimal G protein coupling or in a functional assay designed to test the receptor/G protein hypothesis. In one example, the MCH1 receptor may be hypothesized to inhibit cAMP accumulation through coupling with G alpha sub-units of the G<sub>ai</sub>/G<sub>ao</sub> class. Host cells co-transfected with the MCH1 receptor and appropriate G alpha sub-unit cDNA are stimulated with forskolin +/- MCH1 agonist, as described above in cAMP methods. Intracellular cAMP is extracted for analysis by radioimmunoassay. Other assays may be substituted for cAMP inhibition, including GTPγ<sup>35</sup>S binding assays and inositol phosphate hydrolysis assays. Host cells transfected with MCH1 minus G alpha or with G alpha minus MCH1 would be

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5 tested simultaneously as negative controls. MCH1 receptor expression in transfected cells may be confirmed in radioligand binding studies using membranes from transfected cells. G alpha expression in transfected cells may be confirmed by Western blot analysis of membranes from transfected cells, using antibodies specific for the G protein of interest.

10 The efficiency of the transient transfection procedure is a critical factor for signal to noise in an inhibitory assay, much more so than in a stimulatory assay. If a positive signal present in all cells (such as forskolin-stimulated cAMP accumulation) is inhibited only in the fraction of cells successfully transfected with receptor and G alpha, 15 the signal to noise ratio will be poor. One method for improving the signal to noise ratio is to create a stably transfected cell line in which 100% of the cells express both the receptor and the G alpha subunit. Another method involves transient co-transfection with a third cDNA for a 20 G protein-coupled receptor which positively regulates the signal which is to be inhibited. If the co-transfected cells simultaneously express the stimulatory receptor, the inhibitory receptor, and a requisite G protein for the inhibitory receptor, then a positive signal may be elevated 25 selectively in transfected cells using a receptor-specific agonist. An example involves co-transfection of COS-7 cells with 5-HT4 receptor, MCH1 receptor, and a G alpha sub-unit. Transfected cells are stimulated with a 5-HT4 agonist +/- MCH1 agonist. Cyclic AMP is expected to be elevated only in 30 the cells also expressing MCH1 and the G alpha subunit of interest, and a MCH1-dependent inhibition may be measured with an improved signal to noise ratio.

35 It is to be understood that the cell lines described herein are merely illustrative of the methods used to evaluate the binding and function of the mammalian receptors of the present invention, and that other suitable cells may be used in the assays described herein.

Promiscuous second messenger assays

It is possible to coax receptors of different functional classes to signal through a pre-selected pathway through the use of promiscuous G<sub>o</sub> subunits. For example, by providing a cell based receptor assay system with an exogenously supplied promiscuous G<sub>o</sub> subunit such as G<sub>o16</sub> or a chimeric G<sub>o</sub> subunit such as G<sub>oαq</sub>, a GPCR which normally might prefer to couple through a specific signaling pathway (e.g. G<sub>s</sub>, G<sub>i</sub>, G<sub>q</sub>, G<sub>c</sub>, etc.), can be made to couple through the pathway defined by the promiscuous G<sub>o</sub> subunit and upon agonist activation produce the second messenger associated with that subunit's pathway. In the case of G<sub>o16</sub> and/or G<sub>oαq</sub> this would involve activation of the G<sub>q</sub> pathway and production of the second messenger inositol phosphate. Through similar strategies and tools, it is possible to bias receptor signaling through pathways producing other second messengers such as Ca<sup>++</sup>, cAMP, K<sup>+</sup> currents, etc.

Microphysiometric assay

Because cellular metabolism is intricately involved in and effected by a broad range of cellular events (including receptor activation of various second messenger pathways), the use of microphysiometric measurements of cell metabolism can in principle provide a generic assay of cellular activity arising from the activation of any receptor regardless of the specifics of the receptor's proximal signaling pathway.

General guidelines for cell preparation and microphysiometric recording have been previously reported (Salon, J.A. and Owicki, J.A., 1996). A typical protocol employing transiently transfected CHO cells is as follows; 24 hours prior to recording, transfected cells are harvested and counted. 3 x 10<sup>5</sup> cells are seeded into cell culture capsules (Costar), and allowed to attach to the capsule membrane. 10 hours later (14 hours prior to recording) the

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cell media is switched to serum free F-12 complete to minimize ill-defined metabolic stimulation caused by assorted serum factors.

5 On the day of the experiment the cell capsules are transferred to the microphysiometer (Cytosensor, Molecular Devices Corporation, Sunnyvale, CA) and allowed to equilibrate in recording media (low buffered RPMI 1640, no bicarbonate, no serum) with 0.1% BSA (essentially fatty acid 10 free), during which a baseline measurement of basal metabolic activity is established. The recording paradigm consists of a 100 µl/min flow rate, with a 2 min pump cycle which includes a 30 sec flow interruption during which the rate measurement is taken. Challenges involve a 1 min 20 sec exposure to a drug just prior to the first post challenge rate measurement being taken, followed by two additional pump cycles for a total of 5 min 20 sec drug 15 exposure. Drug is then washed out and rates allowed to return to basal. Reported extracellular acidification rates 20 are expressed as a percentage increase of the peak response over the baseline rate observed just prior to challenge.

#### GPCR ligand library

Functional assays of new receptors such as MCH1 may include 25 a preliminary test of a small library of compounds containing representative agonists for all known GPCRs as well as other compounds which may be agonists for prospective GPCRs or which may be effectors for targets peripherally involved with GPCRs. The collection used in 30 this study comprises approximately 180 compounds (including small molecules, hormones, preprohormones, peptides, etc.) for more than 45 described classes of GPCRs (serotonin, dopamine, noradrenaline, opioids, etc.) and additionally 35 includes ligands for known or suspected but not necessarily pharmacological characterized or cloned GPCR families (such as MCH).

The diversity of the library can be expanded to include

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5                   agonist and antagonist compounds specific for GPCR subtypes, combinatorial peptide and/or small molecule libraries, natural product collections, and the like. To facilitate robotic handling, the substances are distributed as either separate or pooled compound concentrates in 96 well plates and stored frozen as ready to use reagent plates.

Localization of mRNA coding for human MCH1 receptors

10                  Development of probes for MCH1: To facilitate the production of radiolabeled, antisense RNA probes a fragment of the gene encoding rat MCH1 will be subcloned into a plasmid vector containing RNA polymerase promoter sites. The full length cDNA encoding the rat MCH1 will be digested with Pst 1, (nucleotides 905-1194) and this 289 nucleotide fragment will be cloned into the Pst I site of pGEM 3z, containing both sp6 and T7 RNA polymerase promoter sites. The construct will be sequenced to confirm sequence identity and orientation. To synthesize antisense strands of RNA, this construct will be linearized with Hind III or Eco RI (depending on orientation) and T7 or sp6 RNA polymerase will be used to incorporate radiolabeled nucleotide as described below.

25                  A probe coding for the rat glyceraldehyde 3-phosphate dehydrogenase (GAPDH) gene, a constitutively expressed protein, was used concurrently. GAPDH is expressed at a relatively constant level in most tissue and its detection is used to compare expression levels of the rat MCH1 receptors gene in different regions.

30                  Synthesis of probes: MCH1 and GAPDH cDNA sequences preceded by phage polymerase promoter sequences will be used to synthesize radiolabeled riboprobes. Conditions for the synthesis of riboprobes will be: 0.25-1.0 µg linearized DNA plasmid template, 1.5 µl of ATP, GTP, UTP (10 mM each), 3 µl dithiothreitol (0.1 M), 30 units RNAsin RNAse inhibitor, 0.5-1.0 µl (15-20 units/µl) RNA polymerase, 7.0 µl

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transcription buffer (Promega Corp.), and 12.5  $\mu$ l  $\alpha^{32}$ -P-CTP (specific activity 3,000Ci/mmol). 0.1 mM CTP (0.02-1.0  $\mu$ l) will be added to the reactions, and the volume will be adjusted to 35  $\mu$ l with DEPC-treated water. Labeling reactions will be incubated at 37°C for 60 min, after which 5 3 units of RQ1 RNase-free DNase (Promega Corp.) will be added to digest the template. Riboprobes will be separated from unincorporated nucleotides using Microspin S-300 columns (Pharmacia Biotech). TCA precipitation and liquid 10 scintillation spectrometry will be used to measure the amount of label incorporated into the probe. A fraction of all riboprobes synthesized will be size-fractionated on 0.25 mm thick 7M urea, 4.5% acrylamide sequencing gels. These gels will be apposed to storage phosphor screens and the 15 resulting autoradiograph scanned using a phosphorimager (Molecular Dynamics, Sunnyvale, CA) to confirm that the probes synthesized were full-length and not degraded.

**Solution hybridization/ribonuclease protection assay (RPA):**

For solution hybridization 2.0  $\mu$ g of mRNA isolated from 20 tissues will be used. Negative controls consisted of 30  $\mu$ g transfer RNA (tRNA) or no tissue blanks. All mRNA samples will be placed in 1.5-ml microfuge tubes and vacuum dried. Hybridization buffer (40  $\mu$ l of 400 mM NaCl, 20 mM Tris, pH 25 6.4, 2 mM EDTA, in 80% formamide) containing 0.25-2.0 E<sup>6</sup> counts of each probe will be added to each tube. Samples will be heated at 95°C for 15 min, after which the temperature will be lowered to 55°C for hybridization.

After hybridization for 14-18 hr, the RNA/probe mixtures 30 will be digested with RNase A (Sigma) and RNase T1 (Life Technologies). A mixture of 2.0  $\mu$ g RNase A and 1000 units of RNase T1 in a buffer containing 330 mM NaCl, 10 mM Tris (pH 8.0) and 5 mM EDTA (400  $\mu$ l) will be added to each sample 35 and incubated for 90 min at room temperature. After digestion with RNases, 20  $\mu$ l of 10% SDS and 50  $\mu$ g proteinase K will be added to each tube and incubated at 37°C for 15 min. Samples will be extracted with

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phenol/chloroform:isoamyl alcohol and precipitated in 2 volumes of ethanol for 1 hr at -70°C. Pellet Paint (Novagen) will be added to each tube (2.0 µg) as a carrier to facilitate precipitation. Following precipitation, samples will be centrifuged, washed with cold 70% ethanol, and vacuum dried. Samples will be dissolved in formamide loading buffer and size-fractionated on a urea/acrylamide sequencing gel (7.0 M urea, 4.5% acrylamide in Tris-borate-EDTA). Gels will be dried and apposed to storage phosphor screens and scanned using a phosphorimager (Molecular Dynamics, Sunnyvale, CA).

RT-PCR: For the detection of RNA encoding human MCH1, RT-PCR was carried out on mRNA extracted from human tissue. Reverse transcription and PCR reactions were carried out in 50 ml volumes using EZrTth DNA polymerase (Perkin Elmer). Primers with the following sequences were used:

Forward primer (RA SLC1a /MCH F); TCA GCT CGG TTG TGG  
20 GAG CA (SEQ ID NO: 14)  
Reverse primer (RA/SLC1a MCH B); CTT GGA CTT CTT CAC  
GAC (SEQ ID NO: 15)

These primers will amplify a 248 base pair fragment from  
25 nucleotide 169 to 417.

Each reaction contained 0.1 µg mRNA and 0.3µM of each primer. Concentrations of reagents in each reaction were: 300 µM each of GTP; dATP; dCTP; dTTP; 2.5mM Mn(OAc)2; 50 mM Bicine; 115 mM potassium acetate, 8% glycerol and 5 units EZrTth DNA polymerase. All reagents for PCR (except mRNA and oligonucleotide primers) were obtained from Perkin Elmer. Reactions were carried out under the following conditions: 65°C 60 min., 94°C 2 min., (94°C, 1 min., 65°C 1 min) 35 cycles, 72°C 10 min. PCR reactions were size fractionated by gel electrophoresis using 10% polyacrylamide. DNA was stained with SYBR Green I

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(Molecular Probes, Eugene OR) and scanned on a Molecular Dynamics (Sunnyvale CA) Storm 860 in blue fluorescence mode at 450 nM.

- 5 Positive controls for PCR reactions consisted of amplification of the target sequence from a plasmid construct, as well as reverse transcribing and amplifying a known sequence. Negative controls consisted of mRNA blanks, as well as primer and mRNA blanks. To confirm that the mRNA was not contaminated with genomic DNA, samples were digested with RNases before reverse transcription. Integrity of RNA 10 was assessed by amplification of mRNA coding for GAPDH.

Results and Discussion  
Cloning and Sequencing

Discovery of an Expressed Sequence Tag (EST) F07228 in  
5 GENEMBL Homologous to FB41a

A BLAST search of GENEMBL with a Synaptic Pharmaceutical Corporation proprietary sequence, FB41a, resulted in the identification of an EST (accession number F07228) with a high degree of homology to FB41a and somatostatin, opiate 10 and galanin receptors.

Construction and Screening of a Human Hippocampal cDNA  
Library

A human hippocampal cDNA library containing a total of 2.2 15  $\times 10^6$  independent clones with a mean insert size of 3.0 kb was prepared in the expression vector pEXJ.BS. The library was plated on agar plates (ampicillin selection) and glycerol stocks for 450 pools of 5000 independent clones were prepared. Primary glycerol stocks were also grouped 20 together in groups of approximately 10 to create superpools.

Cloning of the full-length sequence of MCH1

Glycerol stocks of the superpools and primary pools from the 25 human hippocampal cDNA library were screened by PCR with F07228 specific primers T579 and T580. One positive primary pool 490, was successively divided into subpools, amplified in LB medium overnight and screened by PCR using primers T579 and T580. One positive subpool, 490-4-10-23 was plated 30 on agar plates (ampicillin selection), and colonies were transferred to nitrocellulose membranes (Schleicher and Schuell, Keene, NH). Filters were hybridized for two days under high stringency conditions with  $10^6$  cpm/ml of a  $^{32}$ P-labeled cDNA probe, T581, designed against the F07228 EST sequence. Filters were washed and apposed to Biomax MS film 35 (Kodak). Seven positive colonies were picked, streaked on LB-AMP plates, and grown overnight. Two individual colonies from each of the original seven were picked and subjected to vector-anchored PCR using the following primer pairs: T95,

T580 and T94, T579. One positive colony, G1, was amplified overnight in TB and processed for plasmid purification. This plasmid was designated TL230 and sequenced on both strands. Nucleotide and peptide sequence analysis were performed with GCG programs (Genetics Computer Group, Madison, WI). A HindIII- KpnI fragment of TL230 was subcloned into the mammalian expression vector pEXJ, and named TL231. The largest open reading frame in this construct contains 1266 nucleotides (Figure 1), which is predicted to encode a protein of 422 amino acids (Figure 2).

There are three in-frame methionines in the amino terminus which could result in a protein of 422, 417 or 353 amino acids. Hydropathy analysis of the protein is consistent with a putative topography of seven transmembrane domains, indicative of the G protein-coupled receptor family (Figure 3). TL231 has been named MCH1.

Database analysis of the sequence of MCH1 revealed that it was most similar to somatostatin receptors. Further database analysis revealed a Genbank submission (accession number AF008650, deposited on October 1, 1997) which appears to be the rat homologue of TL231. AF008650 is 69 nucleotides shorter than MCH1 at the 5'end, and predicts a different initiating methionine. Figures 4 and 5 illustrate the nucleotide and amino acid sequence for the rat MCH1 receptor, respectively.

#### Inositol phosphate response of MCH1-transfected cells

The expression vector (pEXJ) containing the MCH1 cDNA was transfected by electroporation into Cos-7 cells in combination with an expression vector (pEXJ) containing the  $G_{\alpha i}$  subunit. After plating and labeling with [ $^3$ H]-myo-inositol, the transfectants were challenged with a ligand library that included, among other things, melanin concentrating hormone (MCH) (10  $\mu$ M final concentration) and then assayed for inositol phosphate (IP) formation. In five out of the seven screens, cells transfected with MCH1 (with  $G_{\alpha i}$ ) gave an approximately 1.4-fold increase in IP

production as compared to cells transfected with G<sub>q</sub>; alone when challenged with MCH.

Subsequent experiments demonstrated that 10  $\mu$ M MCH was able to stimulate IP release 3.4-fold over basal levels in Cos-7 cells transfected with MCH1 alone, suggesting that this receptor couples through the G<sub>q</sub> signalling pathway. The IP response was shown to be dose-dependent to MCH with an EC<sub>50</sub> value of 9.3  $\pm$  1.7 nM (n=2) and an E<sub>max</sub> of approximately 400% basal (404  $\pm$  72) (Figure 6).

Several additional compounds were tested for their ability to activate MCH1. No dose-responsiveness of inositol phosphate formation could be detected in Cos-7 cells transfected with MCH1 when challenged with somatostatin, haloperidol, or dynorphin A1-13, discounting the possibility that MCH1 encodes a somatostatin-like or opioid-like or sigma-like GPCR subtype (Figure 7)

Microphysiometric response of MCH1-transfected cells to MCH  
CHO cells were transiently transfected with MCH1 using lipofectant, challenged with increasing concentrations of MCH or Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH, and subsequently monitored for changes in extracellular acidification rates. Both ligands produced a dose-dependent increase in acidification rate with an EC<sub>50</sub> value of 8.6 nM for MCH and 51.8 nM for Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH. Neither native CHO cells or mock (pEXJ) transfected CHO cells exhibited a change in acidification rate when exposed to MCH or Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH (Figure 8).

Transcriptional response of MCH1-transfected cells  
Cos-7 cells were transiently transfected with MCH1 and a c-fos- $\beta$ -gal reporter construct by the DEAE-dextran method. The cells were challenged with assorted drugs, including MCH, and transcriptional activity measured by colorimetric assay of  $\beta$ -galactosidase protein expression. Initial single dose challenges with MCH at a concentration of 10  $\mu$ M stimulated c-fos-regulated transcriptional activity approximately 3.9-

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fold over cells challenged with medium only. Cells transfected with only the c-fos- $\beta$ -gal construct showed no response to MCH. Subsequent experimentation showed the transcription activation response to be dose-dependent to MCH with an EC<sub>50</sub> value of 116 nM (Figure 9).

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Binding of [<sup>125</sup>I]Phe<sup>13</sup>,Tyr<sup>19</sup>-MCH in MCH1-transfected cells

Membranes harvested from Cos-7 cells transfected with MCH1 by the DEAE-dextran method exhibited specific binding for [<sup>125</sup>I]Phe<sup>13</sup>-Tyr<sup>19</sup>-MCH (about 80 fmol/mg membrane protein) over mock-transfected cells (about 20 fmol/mg membrane protein) at 0.1 nM radioligand concentration. Specific [<sup>125</sup>I]Phe<sup>13</sup>-Tyr<sup>19</sup>-MCH binding was about 70% of total binding at a radioligand concentration of 0.1 nM (Figure 10).

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Localization of mRNA encoding human MCH1 receptors

RT-PCR was used to assess the presence of MCH1 receptor encoding message in mRNA samples isolated from a variety of human tissues (Table 1, Figure 11). After amplification, PCR reactions were size fractionated on 10% polyacrylamide gels, and stained with SYBR Green I. Images were analyzed using a Molecular Dynamics Storm 860 workstation. The amplified band corresponding to MCH1 receptor (490 base pairs) is indicated (arrow). RT-PCR analysis indicates the distribution of mRNA encoding human MCH1 receptor is widespread throughout all tissues assayed, including both central nervous system tissue and peripheral organs. This widespread distribution implies broad regulatory functions that involve nervous system as well as endocrine mechanisms.

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**Table 1.** Distribution of mRNA coding for human MCH1 receptors.

	<b>Region</b>	<b>human MCH 1</b>	<b>Potential applications</b>
5	<b>liver</b>	+++	Diabetes
	<b>kidney</b>	+++	Hypertension, Electrolyte balance
10	<b>lung</b>	+++	Respiratory disorders, asthma
	<b>heart</b>	+++	Cardiovascular indications
	<b>small intestine</b>	+++	Gastrointestinal disorders
15	<b>striated muscle</b>	+++	Musculoskeletal disorders
	<b>pituitary</b>	+++	Endocrine/neuroendocrine regulation
	<b>whole brain</b>	+++	
	<b>amygdala</b>	+++	Depression, phobias, anxiety, mood disorders
20	<b>cerebral cortex</b>	+++	Sensory and motor integration, cognition
	<b>hippocampus</b>	+++	Cognition/memory
	<b>hypothalamus</b>	+++	appetite/obesity, neuroendocrine regulation
25	<b>spinal cord</b>	+++	Analgesia, sensory modulation and transmission
	<b>cerebellum</b>	+++	Motor coordination
	<b>thalamus</b>	+++	sensory integration
	<b>substantia nigra</b>	+++	Modulation of dopaminergic function. Modulation of motor coordination.
	<b>caudate-putamen</b>	+++	Modulation of dopaminergic function
	<b>fetal brain</b>	+++	Developmental disorders
	<b>fetal lung</b>	+++	Developmental disorders
	<b>fetal kidney</b>	+++	Developmental disorders
	<b>fetal liver</b>	+++	Developmental disorders

The cloning of the gene encoding the human MCH1 receptor has provided the means to explore its physiological role by pharmacological characterization, and by Northern and *in situ* mapping of its mRNA distribution. Further, the 5 availability of the DNA encoding the human MCH1 receptor will facilitate the development of antibodies and antisense technologies useful in defining the functions of the gene products *in vivo*. Antisense oligonucleotides which target mRNA molecules to selectively block translation of the gene 10 products *in vivo* have been used successfully to relate the expression of a single gene with its functional sequelae. Thus, the cloning of this receptor gene provides the means to explore its physiological role in the nervous system and elsewhere, and may thereby help to elucidate 15 structure/function relationships within the GPCR superfamily.

The presence of three different potential starting codons in the cDNA sequence of TL231 opens the question of which of the possible transcripts yields an active MCH receptor. In 20 order to establish whether a transcript of the first and second starting codons of TL231 encode a functional human MCH receptor, methionines 6 and 70 of TL231 were mutated to alanine (construct R114; See Figure 12). The third methionine at position 70 was also mutated to an alanine 25 (construct R106; See Figure 12). Transfections of TL231, R106 or R114 into COS-7 cells all resulted in MCH-mediated increases of intracellular calcium, as measured by a fluorescent intensity plate reader in cells loaded with the calcium dye fluo-3 (FLIPR, Molecular Devices). As shown in 30 Table 2, COS-7 cells transfected with TL231, R106, R114 and B0120 showed dose-related mobilization of intracellular calcium when exposed to increasing concentrations of MCH with similar maximal responses and EC<sub>50</sub> values. These data demonstrate that transcripts starting at the first and/or 35 second and third methionine of TL231 encode a functional

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human MCH receptor.

Table 2.

Transfected Construct	Response to Melanin Concentrating Hormone*	
	EC50 (nM)	Max. Response (RFU**)
TL231	60, 12	3,535, 14,000
R114	98, 9	2,267, 1,550
R106	85, 55	4642, 2000
BO120	12, 3.5	30,000, 25,000

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\*Results from two independent experiments

\*\* RFU = relative fluorescence units

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What is claimed is:

1. An isolated nucleic acid encoding a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.  
5
2. The nucleic acid of claim 1, wherein the nucleic acid is DNA.  
  
3. The DNA of claim 2, wherein the DNA is cDNA.  
10
4. The DNA of claim 2, wherein the DNA is genomic DNA.  
  
5. The nucleic acid of claim 1, wherein the nucleic acid is RNA.  
15
6. The nucleic acid of claim 1, wherein the human MCH1 receptor has an amino acid sequence identical to that encoded by the plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197).  
20
7. The nucleic acid of claim 1, wherein the human MCH1 receptor comprises an amino acid sequence as shown in Figure 2 (SEQ ID NO: 2).  
  
25
8. The nucleic acid of claim 1, wherein the mutant human MCH1 receptor comprises an amino acid sequence as shown in Figure 13 (SEQ ID NO: 27).

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9. The nucleic acid of claim 1, wherein the mutant human MCH1 receptor comprises an amino acid sequence as shown in Figure 14 (SEQ ID NO: 28).  
5       10. The nucleic acid of claim 1, wherein the mutant human MCH1 receptor comprises an amino acid sequence as shown in Figure 15 (SEQ ID NO: 29).  
  
11. A purified human MCH1 receptor protein.  
10  
12. A vector comprising the nucleic acid of claim 1.  
  
13. The vector of claim 12 adapted for expression in a cell which comprises the regulatory elements necessary for expression of the nucleic acid in the cell operatively linked to the nucleic acid encoding the receptor so as to permit expression thereof, wherein the cell is a bacterial, amphibian, yeast, insect or mammalian cell.  
15  
20       14. The vector of claim 13, wherein the vector is a baculovirus.  
  
15. The vector of claim 12, wherein the vector is a plasmid.  
  
25       16. The plasmid of claim 15 designated pEXJ.HR-TL231 (ATCC Accession No. 203197).  
  
17. A cell comprising the vector of claim 13.

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18. A cell of claim 17, wherein the cell is a non-mammalian cell.

19. A cell of claim 18, wherein the non-mammalian cell is a Xenopus oocyte cell or a Xenopus melanophore cell.

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20. A cell of claim 17, wherein the cell is a mammalian cell.

10 21. A mammalian cell of claim 20, wherein the cell is a COS-7 cell, a 293 human embryonic kidney cell, a NIH-3T3 cell, a LM(tk-) cell, a mouse Y1 cell, or a CHO cell.

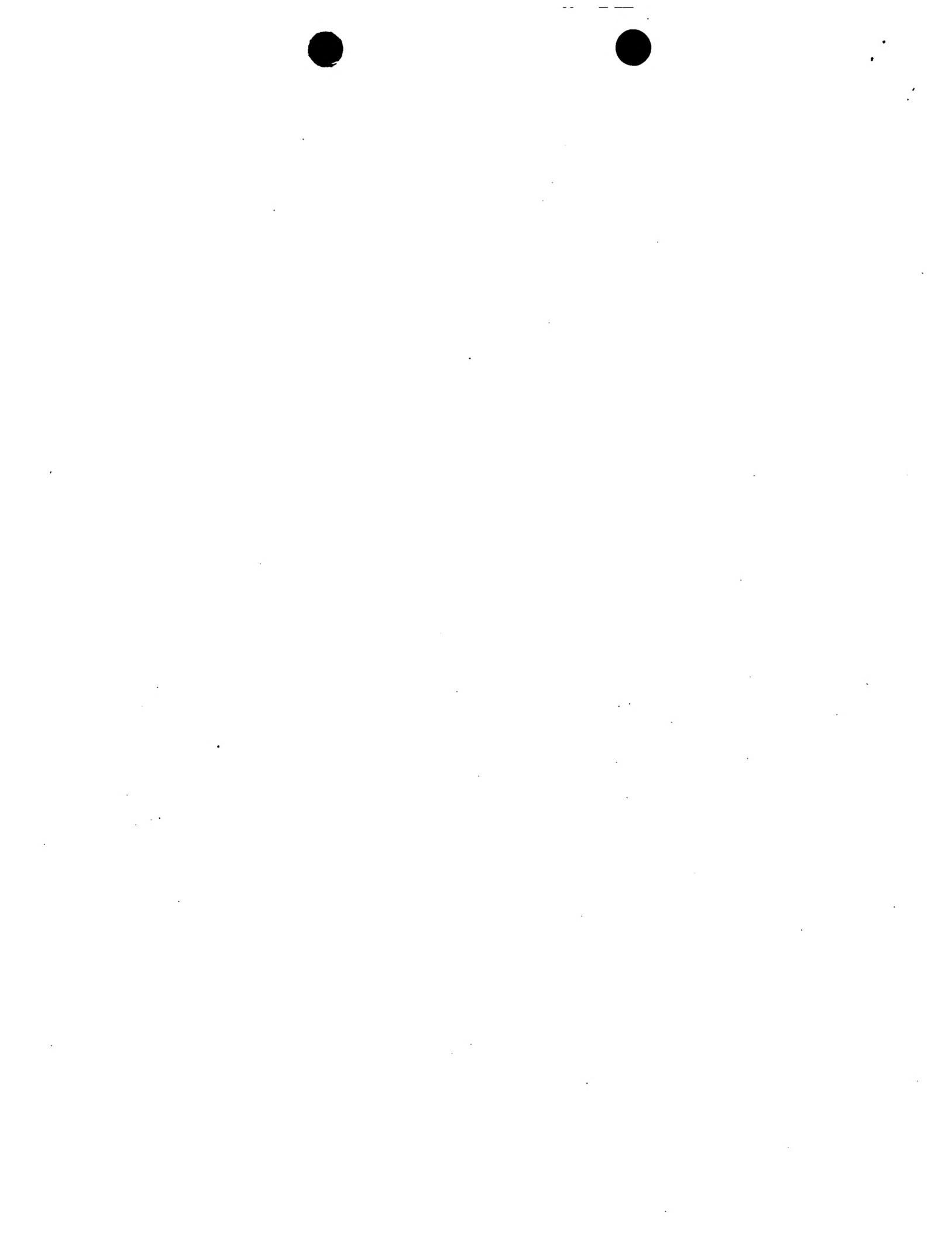
22. An insect cell comprising the vector of claim 13.

15

23. An insect cell of claim 22, wherein the insect cell is an Sf9 cell, an Sf21 cell or a Trichoplusia ni 5B-4 cell.

20 24. A membrane preparation isolated from the cell of claim 17.

25 25. A nucleic acid probe comprising at least 15 nucleotides which specifically hybridizes with a nucleic acid encoding a human MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within one of the two strands of the nucleic acid encoding a human MCH1 receptor present in plasmid pEXJ.HR-T231 (ATCC Accession No. 203197).





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- 5           26. A nucleic acid probe comprising at least 15 nucleotides which specifically hybridizes with a nucleic acid encoding a human MCH1 receptor, wherein the probe has a unique sequence corresponding to a sequence present within (a) the nucleic acid sequence shown in Figure 1 (SEQ ID NO: 1) or (b) the reverse complement thereof.
- 10           27. The nucleic acid probe of claim 25 or 26, wherein the nucleic acid is DNA.
- 15           28. The nucleic acid probe of claim 25 or 26, wherein the nucleic acid is RNA.
- 20           29. An antisense oligonucleotide having a sequence capable of specifically hybridizing to the RNA of claim 5, so as to prevent translation of the RNA.
- 25           30. An antisense oligonucleotide having a sequence capable of specifically hybridizing to the genomic DNA of claim 4.
- 30           31. An antisense oligonucleotide of claim 29 or 30, wherein the oligonucleotide comprises chemically modified nucleotides or nucleotide analogues.
- 35           32. An antibody capable of binding to a human MCH1 receptor encoded by the nucleic acid of claim 1.
- 40           33. An agent capable of competitively inhibiting the binding of the antibody of claim 32 to a human MCH1 receptor.

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34. An antibody of claim 32, wherein the antibody is a monoclonal antibody or antisera.
- 5       35. A pharmaceutical composition comprising (a) an amount of the oligonucleotide of claim 29 capable of passing through a cell membrane and effective to reduce expression of a human MCH1 receptor and (b) a pharmaceutically acceptable carrier capable of passing through the cell membrane.
- 10      36. A pharmaceutical composition of claim 35, wherein the oligonucleotide is coupled to a substance which inactivates mRNA.
- 15      37. A pharmaceutical composition of claim 36, wherein the substance which inactivates mRNA is a ribozyme.
- 20      38. A pharmaceutical composition of claim 35, wherein the pharmaceutically acceptable carrier comprises a structure which binds to a human MCH1 receptor on a cell capable of being taken up by the cells after binding to the structure.
- 25      39. A pharmaceutical composition of claim 35, wherein the pharmaceutically acceptable carrier is capable of binding to a human MCH1 receptor which is specific for a selected cell type.
- 30      40. A pharmaceutical composition which comprises an amount of the antibody of claim 32 effective to block binding of a ligand to a human MCH1 receptor and a

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pharmaceutically acceptable carrier.

41. A transgenic, nonhuman mammal expressing DNA encoding a human MCH1 receptor of claim 1.

5

42. A transgenic, nonhuman mammal comprising a homologous recombination knockout of the native human MCH1 receptor.

10

43. A transgenic, nonhuman mammal whose genome comprises antisense DNA complementary to the DNA encoding a human MCH1 receptor of claim 1 so placed within the genome as to be transcribed into antisense mRNA which is complementary to mRNA encoding the human MCH1 receptor and which hybridizes to mRNA encoding the human MCH1 receptor, thereby reducing its translation.

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44. The transgenic, nonhuman mammal of claim 41 or 42, wherein the DNA encoding the human MCH1 receptor additionally comprises an inducible promoter.

20

45. The transgenic, nonhuman mammal of claim 41 or 42, wherein the DNA encoding the human MCH1 receptor additionally comprises tissue specific regulatory elements.

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46. A transgenic, nonhuman mammal of claim 41, 42, or 43, wherein the transgenic, nonhuman mammal is a mouse.

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47. A process for identifying a chemical compound which

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5 specifically binds to a mammalian MCH1 receptor which comprises contacting cells comprising DNA encoding, and expressing on their cell surface, the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO 10 cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

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20 48. A process for identifying a chemical compound which specifically binds to a mammalian MCH1 receptor which comprises contacting a membrane preparation from cells comprising DNA encoding, and expressing on their cell surface, the mammalian MCH1 receptor, with the compound under conditions suitable for binding, and detecting specific binding of the chemical compound to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to 25 a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

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49. The process of claim 47 or 48, wherein the mammalian MCH1 receptor is a human MCH1 receptor.
- 5 50. The process of claim 47 or 48, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.
- 10 51. The process of claim 47 or 48, wherein the mammalian MCH1 receptor has substantially the same amino acid sequence as the sequence of the human MCH1 receptor encoded by plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197).
- 15 52. The process of claim 47 or 48, wherein the mammalian MCH1 receptor comprises substantially the same amino acid sequence as that shown in Figure 2 (SEQ ID NO: 2).
- 20 53. The process of claim 47 or 48, wherein the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 2 (SEQ ID NO: 2).
- 25 54. The process of claim 47 or 48, wherein the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 13 (SEQ ID NO: 27).
55. The process of claim 47 or 48, wherein the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 14 (SEQ ID NO: 28).

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56. The process of claim 47 or 48, wherein the mammalian MCH1 receptor comprises the amino acid sequence shown in Figure 15 (SEQ ID NO: 29).
- 5 57. The process of claim 47 or 48, wherein the compound is not previously known to bind to a mammalian MCH1 receptor.
58. A compound identified by the process of claim 57.
- 10 59. A process of claim 47 or 48, wherein the cell is an insect cell.
- 15 60. The process of claim 47 or 48, wherein the cell is a mammalian cell.
61. The process of claim 60, wherein the cell is nonneuronal in origin.
- 20 62. The process of claim 61, wherein the nonneuronal cell is a COS-7 cell, 293 human embryonic kidney cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.
- 25 63. A process of claim 60, wherein the compound is a compound not previously known to bind to a mammalian MCH1 receptor.

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64. A compound identified by the process of claim 63.

65. A process involving competitive binding for identifying  
5 a chemical compound which specifically binds to a  
mammalian MCH1 receptor which comprises contacting cells  
expressing on their cell surface the mammalian MCH1  
receptor, with both the chemical compound and a second  
10 chemical compound known to bind to the receptor, and  
separately with only the second chemical compound, under  
conditions suitable for binding of both compounds, and  
detecting specific binding of the chemical compound to  
the mammalian MCH1 receptor, a decrease in the binding  
15 of the second chemical compound to the mammalian MCH1  
receptor in the presence of the chemical compound  
indicating that the chemical compound binds to the  
mammalian MCH1 receptor, wherein the cells do not  
normally express the mammalian MCH1 receptor and the DNA  
20 encoding the mammalian MCH1 receptor (a) hybridizes to  
a nucleic acid having the defined sequence shown in  
Figure 1 (SEQ ID NO: 1) under low stringency conditions  
or a sequence complementary thereto and (b) is further  
characterized by its ability to cause a change in the pH  
25 of a culture of CHO cells when a MCH1 ligand is added to  
the culture and the CHO cells contain the nucleic acid  
which hybridized to the nucleic acid having the defined  
sequence or its complement.

66. A process involving competitive binding for identifying  
30 a chemical compound which specifically binds to a  
mammalian MCH1 receptor which comprises contacting a  
membrane preparation from cells expressing on their cell  
surface the mammalian MCH1 receptor, with both the  
chemical compound and a second chemical compound known  
35 to bind to the receptor, and separately with only the  
second chemical compound, under conditions suitable for  
binding of both compounds, and detecting specific

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binding of the chemical compound to the mammalian MCH1 receptor, a decrease in the binding of the second chemical compound to the mammalian MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the mammalian MCH1 receptor, wherein the cells do not normally express the mammalian MCH1 receptor and the DNA encoding the mammalian MCH1 receptor (a) hybridizes to a nucleic acid having the defined sequence shown in Figure 1 (SEQ ID NO: 1) under low stringency conditions or a sequence complementary thereto and (b) is further characterized by its ability to cause a change in the pH of a culture of CHO cells when a MCH1 ligand is added to the culture and the CHO cells contain the nucleic acid which hybridized to the nucleic acid having the defined sequence or its complement.

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67. A process of claim 65 or 66, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.
68. A process of claim 65 or 66, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.
69. The process of claim 65 or 66, wherein the cell is an insect cell.
70. The process of claim 65 or 66, wherein the cell is a mammalian cell.
71. The process of claim 70, wherein the cell is nonneuronal

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in origin.

5           72. The process of claim 71, wherein the nonneuronal cell is a COS-7 cell, 293 human embryonic kidney cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.

10           73. The process of claim 70, wherein the compound is not previously known to bind to a mammalian MCH1 receptor.

15           74. A compound identified by the process of claim 73.

20           75. A method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises

25           (a)       contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor, under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor;

30           (b)       determining whether the binding of a compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so

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(c) separately determining the binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.

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76. A method of screening a plurality of chemical compounds not known to bind to a mammalian MCH1 receptor to identify a compound which specifically binds to the mammalian MCH1 receptor, which comprises

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(a) contacting a membrane preparation from cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the plurality of compounds not known to bind specifically to the mammalian MCH1 receptor under conditions permitting binding of compounds known to bind the mammalian MCH1 receptor;

20

(b) determining whether the binding of a compound known to bind to the mammalian MCH1 receptor is reduced in the presence of the compounds within the plurality of compounds, relative to the binding of the compound in the absence of the plurality of compounds; and if so

25

(c) separately determining the binding to the mammalian MCH1 receptor of compounds included in the plurality of compounds, so as to thereby identify the compound which specifically binds to the mammalian MCH1 receptor.

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77. A method of claim 75 or 76, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

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78. A method of claim 75 or 76, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.

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79. A method of claim 75 or 76, wherein the cell is a mammalian cell.

80. A method of claim 79, wherein the mammalian cell is non-neuronal in origin.

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81. The method of claim 80, wherein the non-neuronal cell is a COS-7 cell, a 293 human embryonic kidney cell, a LM(tk-) cell, a CHO cell, a mouse Y1 cell, or an NIH-3T3 cell.

20  
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82. A method of detecting expression of a mammalian MCH1 receptor by detecting the presence of mRNA coding for the mammalian MCH1 receptor which comprises obtaining total mRNA from the cell and contacting the mRNA so obtained with the nucleic acid probe of any of claims 25, 26, 27, or 28 under hybridizing conditions, detecting the presence of mRNA hybridizing to the probe, and thereby detecting the expression of the mammalian MCH1 receptor by the cell.

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83. A method of detecting the presence of a mammalian MCH1 receptor on the surface of a cell which comprises

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5 contacting the cell with the antibody of claim 32 under conditions permitting binding of the antibody to the receptor, detecting the presence of the antibody bound to the cell, and thereby detecting the presence of the mammalian MCH1 receptor on the surface of the cell.

10 84. A method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a transgenic, nonhuman mammal of claim 44 whose levels of human MCH1 receptor activity are varied by use of an inducible promoter which regulates human MCH1 receptor expression.

15 85. A method of determining the physiological effects of varying levels of activity of human MCH1 receptors which comprises producing a panel of transgenic, nonhuman mammals of claim 44, each expressing a different amount of human MCH1 receptor.

20 86. A method for identifying an antagonist capable of alleviating an abnormality, wherein the abnormality is alleviated by decreasing the activity of a human MCH1 receptor comprising administering a compound to the transgenic, nonhuman mammal of claim 41, 44, 45, or 46, and determining whether the compound alleviates the physical and behavioral abnormalities displayed by the transgenic, nonhuman mammal as a result of overactivity of a human MCH1 receptor, the alleviation of the abnormality identifying the compound as an antagonist.

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87. An antagonist identified by the method of claim 86.

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88. A pharmaceutical composition comprising an antagonist of  
claim 87 and a pharmaceutically acceptable carrier.
- 5                   89. A method of treating an abnormality in a subject wherein  
the abnormality is alleviated by decreasing the activity  
of a human MCH1 receptor which comprises administering  
to the subject an effective amount of the pharmaceutical  
composition of claim 88, thereby treating the  
abnormality.
- 10                  90. A method for identifying an agonist capable of  
alleviating an abnormality in a subject wherein the  
abnormality is alleviated by increasing the activity of  
a human MCH1 receptor comprising administering a  
compound to the transgenic, nonhuman mammal of claim 41,  
44, 45, or 46, and determining whether the compound  
alleviates the physical and behavioral abnormalities  
displayed by the transgenic, nonhuman mammal, the  
alleviation of the abnormality identifying the compound  
as an agonist.
- 15                  91. An agonist identified by the method of claim 90.
- 20                  92. A pharmaceutical composition comprising an agonist of  
claim 91 and a pharmaceutically acceptable carrier.
- 25                  93. A method of treating an abnormality in a subject wherein  
the abnormality is alleviated by increasing the  
activity of a human MCH1 receptor which comprises  
administering to the subject an effective amount of the  
pharmaceutical composition of claim 92, thereby treating  
the abnormality.

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94. A method for diagnosing a predisposition to a disorder associated with the activity of a specific mammalian allele which comprises:

5 (a) obtaining DNA of subjects suffering from the disorder;

10 (b) performing a restriction digest of the DNA with a panel of restriction enzymes;

(c) electrophoretically separating the resulting DNA fragments on a sizing gel;

15 (d) contacting the resulting gel with a nucleic acid probe capable of specifically hybridizing with a unique sequence included within the sequence of a nucleic acid molecule encoding a human MCH1 receptor and labeled with a detectable marker;

20 (e) detecting labeled bands which have hybridized to the DNA encoding a human MCH1 receptor of claim 1 labeled with a detectable marker to create a unique band pattern specific to the DNA of subjects suffering from the disorder;

25 (f) preparing DNA obtained for diagnosis by steps (a) - (e); and

30 (g) comparing the unique band pattern specific to

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the DNA of subjects suffering from the disorder from step (e) and the DNA obtained for diagnosis from step (f) to determine whether the patterns are the same or different and to diagnose thereby predisposition to the disorder if the patterns are the same.

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95. The method of claim 94, wherein a disorder associated with the activity of a specific mammalian allele is diagnosed.

96. A method of preparing the purified human MCH1 receptor of claim 11 which comprises:

15

(a) inducing cells to express the human MCH1 receptor;

(b) recovering the human MCH1 receptor from the induced cells; and

20

(c) purifying the human MCH1 receptor so recovered.

97. A method of preparing the purified human MCH1 receptor of claim 11 which comprises:

25

(a) inserting nucleic acid encoding the human MCH1 receptor in a suitable vector;

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- (b) introducing the resulting vector in a suitable host cell;
- 5 (c) placing the resulting cell in suitable condition permitting the production of the isolated human MCH1 receptor;
- 10 (d) recovering the human MCH1 receptor produced by the resulting cell; and
- (e) purifying the human MCH1 receptor so recovered.
- 15 98. A process for determining whether a chemical compound is a mammalian MCH1 receptor agonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound under conditions permitting the activation of the mammalian MCH1 receptor, and detecting an increase in mammalian MCH1 receptor activity, so as to thereby determine whether the compound is a mammalian MCH1 receptor agonist.
- 20 99. A process for determining whether a chemical compound is a mammalian MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the mammalian MCH1 receptor with the compound in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting the activation of the mammalian MCH1 receptor, and detecting a decrease in mammalian MCH1 receptor activity, so as to thereby determine whether the compound is a mammalian MCH1 receptor antagonist.
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- 30

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100. A process of claim 98 or 99, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.
- 5
101. A process of claim 98 or 99, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.
102. A pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor agonist determined by the process of claim 98 effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.
- 15 103. A pharmaceutical composition of claim 102, wherein the mammalian MCH1 receptor agonist is not previously known.
- 20 104. A pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor antagonist determined by the process of claim 99 effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.
- 25 105. A pharmaceutical composition of claim 104, wherein the mammalian MCH1 receptor antagonist is not previously known.
- 30 106. A process for determining whether a chemical compound specifically binds to and activates a mammalian MCH1 receptor, which comprises contacting

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5                   cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with the chemical compound under conditions suitable for activation of the mammalian MCH1 receptor, and measuring the second messenger response in the presence and in the absence of the chemical compound, a change in the second messenger response in the presence of the chemical compound indicating that the compound activates the mammalian MCH1 receptor.

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107.               The process of claim 106, wherein the second messenger response comprises chloride channel activation and the change in second messenger is an increase in the level of inward chloride current.

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108.              A process for determining whether a chemical compound specifically binds to and inhibits activation of a mammalian MCH1 receptor, which comprises separately contacting cells producing a second messenger response and expressing on their cell surface the mammalian MCH1 receptor, wherein such cells do not normally express the mammalian MCH1 receptor, with both the chemical compound and a second chemical compound known to activate the mammalian MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the mammalian MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second

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chemical compound indicating that the chemical compound inhibits activation of the mammalian MCH1 receptor.

- 5        109. The process of claim 108, wherein the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.
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- 15        110. A process of any of claims 106, 107, 108, or 109, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.
- 20        111. A process of any of claims 106, 107, 108, or 109, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.
- 25        112. The process of any of claims 106, 107, 108, 109, or 110, wherein the cell is an insect cell.
- 30        113. The process of any of claims 106, 107, 108, 109, or 110, wherein the cell is a mammalian cell.
- 30        114. The process of claim 113, wherein the mammalian cell is nonneuronal in origin.

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115. The process of claim 114, wherein the nonneuronal cell is a COS-7 cell, CHO cell, 293 human embryonic kidney cell, NIH-3T3 cell or LM(tk-) cell.
- 5 116. The process of claim 106, 107, 108, or 109, wherein the compound is not previously known to bind to a mammalian MCH1 receptor.
- 10 117. A compound determined by the process of claim 116.
118. A pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor agonist determined by the process of claim 106 or 107 effective to increase activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.
- 15 119. A pharmaceutical composition of claim 118, wherein the mammalian MCH1 receptor agonist is not previously known.
- 20 120. A pharmaceutical composition which comprises an amount of a mammalian MCH1 receptor antagonist determined by the process of claim 108 or 109 effective to reduce activity of a mammalian MCH1 receptor and a pharmaceutically acceptable carrier.
- 25 121. A pharmaceutical composition of claim 120, wherein the mammalian MCH1 receptor antagonist is not previously known.

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122. A method of screening a plurality of chemical compounds not known to activate a mammalian MCH1 receptor to identify a compound which activates the mammalian MCH1 receptor which comprises:

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(a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds not known to activate the mammalian MCH1 receptor, under conditions permitting activation of the mammalian MCH1 receptor;

(b) determining whether the activity of the mammalian MCH1 receptor is increased in the presence of the compounds; and if so

(c) separately determining whether the activation of the mammalian MCH1 receptor is increased by each compound included in the plurality of compounds, so as to thereby identify the compound which activates the mammalian MCH1 receptor.

123. A method of claim 122, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

124. A method of claim 122, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.

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125. A method of screening a plurality of chemical compounds not known to inhibit the activation of a mammalian MCH1 receptor to identify a compound which inhibits the activation of the mammalian MCH1 receptor, which comprises:

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(a) contacting cells transfected with and expressing the mammalian MCH1 receptor with the plurality of compounds in the presence of a known mammalian MCH1 receptor agonist, under conditions permitting activation of the mammalian MCH1 receptor;

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(b) determining whether the activation of the mammalian MCH1 receptor is reduced in the presence of the plurality of compounds, relative to the activation of the mammalian MCH1 receptor in the absence of the plurality of compounds; and if so

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(c) separately determining the inhibition of activation of the mammalian MCH1 receptor for each compound included in the plurality of compounds, so as to thereby identify the compound which inhibits the activation of the mammalian MCH1 receptor.

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126. A method of claim 125, wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

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127. A method of claim 125, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.
- 5        128. A method of any of claims 123, 124, 125, 126, or 127, wherein the cell is a mammalian cell.
- 10      130. The method of claim 129, wherein the non-neuronal cell is a COS-7 cell, a 293 human embryonic kidney cell, a LM(tk-) cell or an NIH-3T3 cell.
- 15      131. A pharmaceutical composition comprising a compound identified by the method of claim 123 or 124 effective to increase mammalian MCH1 receptor activity and a pharmaceutically acceptable carrier.
- 20      132. A pharmaceutical composition comprising a compound identified by the method of claim 125 or 126 effective to decrease mammalian MCH1 receptor activity and a pharmaceutically acceptable carrier.
- 25      133. A method of treating an abnormality in a subject wherein the abnormality is alleviated by increasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor agonist effective to treat the abnormality.

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134. A method of claim 133, wherein the abnormality is a regulation of a steroid or pituitary hormone disorder, an epinephrine release disorder, a gastrointestinal disorder, a cardiovascular disorder, an electrolyte balance disorder, hypertension, diabetes, a respiratory disorder, asthma, a reproductive function disorder, an immune disorder, an endocrine disorder, a musculoskeletal disorder, a neuroendocrine disorder, a cognitive disorder, a memory disorder, a sensory modulation and transmission disorder, a motor coordination disorder, a sensory integration disorder, a motor integration disorder, a dopaminergic function disorder, a sensory transmission disorder, an olfaction disorder, a sympathetic innervation disorder, pain, psychotic behavior, morphine tolerance, opiate addiction, an affective disorder, a stress-related disorder, a fluid-balance disorder, a seizure disorder, or migraine.
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135. A method of treating an abnormality in a subject wherein the abnormality is alleviated by decreasing the activity of a mammalian MCH1 receptor which comprises administering to the subject an amount of a compound which is a mammalian MCH1 receptor antagonist effective to treat the abnormality.
- 30
136. A method of claim 135, wherein the abnormality is a regulation of a steroid or pituitary hormone disorder, an epinephrine release disorder, a gastrointestinal disorder, a cardiovascular disorder, an electrolyte balance disorder, hypertension, diabetes, a respiratory disorder, asthma, a reproductive function disorder, an immune disorder, an endocrine disorder, a musculoskeletal disorder, a neuroendocrine disorder, a cognitive
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disorder, a memory disorder, a sensory modulation  
and transmission disorder, a motor coordination  
disorder, a sensory integration disorder, a motor  
integration disorder, a dopaminergic function  
disorder, a sensory transmission disorder, an  
olfaction disorder, a sympathetic innervation  
disorder, pain, psychotic behavior, morphine  
tolerance, opiate addiction, an affective disorder,  
a stress-related disorder, a fluid-balance disorder,  
a seizure disorder, or migraine.

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137. A process for making a composition of matter which  
specifically binds to a mammalian MCH1 receptor  
which comprises identifying a chemical compound  
using the process of any of claims 47, 48, 65, 66,  
75, or 76 and then synthesizing the chemical  
compound or a novel structural and functional analog  
or homolog thereof.

138. A process for making a composition of matter which  
specifically binds to a mammalian MCH1 receptor  
which comprises identifying a chemical compound  
using the process of any of claims 98, 106, or 122  
and then synthesizing the chemical compound or a  
novel structural and functional analog or homolog  
thereof.

139. A process for making a composition of matter which  
specifically binds to a mammalian MCH1 receptor  
which comprises identifying a chemical compound  
using the process of any of claims 99, 108, or 125  
and then synthesizing the chemical compound or a  
novel structural and functional analog or homolog  
thereof.

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140. The process of any of claims 137, 138, or 139,  
wherein the mammalian MCH1 receptor is a human MCH1  
receptor or a mutant of such human MCH1 receptor  
which is activated by MCH or an analog or homolog  
thereof.
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141. The process of any of claims 137, 138, or 139,  
wherein the mammalian MCH1 receptor is a human MCH1  
receptor.
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142. A process for preparing a composition which  
comprises admixing a pharmaceutically acceptable  
carrier and a therapeutically effective amount of a  
chemical compound identified by the process of any  
15 of claims 47, 48, 65, 66, 75, or 76 or a novel  
structural and functional analog or homolog thereof.
143. A process for preparing a composition which  
comprises admixing a pharmaceutically acceptable  
carrier and a therapeutically effective amount of a  
chemical compound identified by the process of any  
20 of claims 98, 106, or 122 or a novel structural and  
functional analog or homolog thereof.
- 25 144. A process for preparing a composition which  
comprises admixing a pharmaceutically acceptable  
carrier and a therapeutically effective amount of a  
chemical compound identified by the process of any  
of claims 99, 108, or 125 or a novel structural and  
functional analog or homolog thereof.
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145. The process of any of claims 142, 143, or 144,

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wherein the mammalian MCH1 receptor is a human MCH1 receptor or a mutant of such human MCH1 receptor which is activated by MCH or an analog or homolog thereof.

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146. The process of any of claims 142, 143, or 144, wherein the mammalian MCH1 receptor is a rat MCH1 receptor.

10        147. A process for determining whether a chemical compound is a human MCH1 receptor antagonist which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the DNA encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH, and the cells do not express the MCH1 receptor prior to transfecting them.

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148. A process for determining whether a chemical compound specifically binds to and inhibits activation of a human MCH1 receptor, which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the DNA

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encoding the human MCH1 receptor comprises the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.

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149. The process of claim 148, wherein the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

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30 150. A method of screening a plurality of chemical compounds not known to inhibit the activation of a human MCH1 receptor to identify a compound which inhibits the activation of the human MCH1 receptor, which comprises:

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153. The process of any of claims 147, 148 or 150, wherein the cell is a mammalian cell which is nonneuronal in origin.

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154. The process of any of claims 147, 148 or 150, wherein the cell is a COS-7 cell, a CHO cell, a 293 human embryonic kidney cell, a NIH-3T3 cell, a mouse Y1 cell, or a LM(tk-) cell.

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155. A process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession

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No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

156. A process for making a composition of matter which specifically binds to a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting a membrane preparation from cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

157. A process for making a composition of matter which is a human MCH1 receptor antagonist which comprises identifying a chemical compound which is a human MCH1 receptor antagonist and then synthesizing the

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5           chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as a human MCH1 receptor antagonist by a process which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

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20         158. A process for making a composition of matter which specifically binds to and inhibits the activation of a human MCH1 receptor which comprises identifying a chemical compound which specifically binds to and inhibits the activation of the human MCH1 receptor and then synthesizing the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to and inhibiting the activation of the human MCH1 receptor by a process which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC

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Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.

159. The process of claim 158, wherein the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

160. A process for preparing a composition which comprises identifying a chemical compound which specifically binds to a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process involving competitive binding which comprises contacting cells expressing on their cell surface the human MCH1 receptor, with both the chemical

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5 compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession 10 No. 203197), and the second chemical compound is MCH 15 or a homolog or analog of MCH.

161. A process for preparing a composition which 20 comprises identifying a chemical compound which specifically binds to a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as binding to the human MCH1 receptor by a process 25 involving competitive binding which comprises contacting a membrane preparation from cells expressing on their cell surface the human MCH1 receptor, with both the chemical compound and a second chemical compound known to bind to the receptor, and separately with only the second chemical compound, under conditions suitable for binding of both compounds, and detecting the extent 30 of specific binding of the chemical compound to the human MCH1 receptor, a decrease in the binding of the second chemical compound to the human MCH1 receptor in the presence of the chemical compound indicating that the chemical compound binds to the 35

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human MCH1 receptor, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the second chemical compound is MCH or a homolog or analog of MCH.

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162. A process for preparing a composition which comprises identifying a chemical compound which is a human MCH1 receptor antagonist, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the chemical compound is identified as a human MCH1 receptor antagonist by a process which comprises contacting cells transfected with and expressing DNA encoding the human MCH1 receptor with the compound in the presence of a known human MCH1 receptor agonist, under conditions permitting the activation of the human MCH1 receptor, and detecting a decrease in human MCH1 receptor activity, so as to thereby determine whether the compound is a human MCH1 receptor antagonist, wherein the cells do not normally express the human MCH1 receptor, the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), and the known human MCH1 receptor agonist is MCH or a homolog or analog of MCH.

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163. A process for preparing a composition which comprises identifying a chemical compound which specifically binds to and inhibits the activation of a human MCH1 receptor, and then admixing a carrier and the chemical compound or a structural and functional analog or homolog thereof, wherein the

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chemical compound is identified as binding to and inhibiting activation of the human MCH1 receptor by a process which comprises separately contacting cells expressing on their cell surface the human MCH1 receptor and producing a second messenger response upon activation of the human MCH1 receptor, wherein such cells do not normally express the human MCH1 receptor and the human MCH1 receptor is encoded by nucleic acid comprising the sequence shown in Figure 1 (Seq. ID No. 1) or contained in plasmid pEXJ.HR-TL231 (ATCC Accession No. 203197), with both the chemical compound and a second chemical compound known to activate the human MCH1 receptor, and with only the second chemical compound, under conditions suitable for activation of the human MCH1 receptor, and measuring the second messenger response in the presence of only the second chemical compound and in the presence of both the second chemical compound and the chemical compound, a smaller change in the second messenger response in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound indicating that the chemical compound inhibits activation of the human MCH1 receptor, wherein the second chemical compound is MCH or a homolog or analog of MCH.

164. The process of claim 163, wherein the second messenger response comprises chloride channel activation and the change in second messenger response is a smaller increase in the level of inward chloride current in the presence of both the chemical compound and the second chemical compound than in the presence of only the second chemical compound.

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165. The process of any of claims 155, 156, 157, 158,  
160, 161, 162, or 163, wherein the cell is an insect  
cell.

5 166. The process of any of claims 155, 156, 157, 158,  
160, 161, 162, or 163, wherein the cell is a  
mammalian cell.

10 167. The process of claim 166, wherein the mammalian cell  
is nonneuronal in origin.

15 168. The process of claim 167, wherein the nonneuronal  
cell is a COS-7 cell, a 293 human embryonic kidney  
cell, a CHO cell, a NIH-3T3 cell, a mouse Y1 cell,  
or a LM(tk-) cell.

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FIGURE 1

1 ATGTCAGTGGGAGCCATGAAGAAGGGAGTGGGAGGGCAGTTGGCTTGGAGGGGGAGC  
 61 GGCTGCCAGGGCTACGGAGGAAGACCCCTTCCCGACTGGGGCTGGCTGGCTCCGGGACAA  
 121 GGTGGCAGGGCTGGAGGCTGGCGAGCCCTGGGTGGAGGGAGCTCAGCTCGGTG  
 181 TGGAAGCAGGGGACCCGCACTGGCTGGATGGACCTGGAGCCCTGGAAAGCTGGT  
 241 CCAATGCCAGCAACACCTCTGATGGCCCCGATAACCTCACTCATGCCTTCGGGACCC  
 301 CGCACGGGGAGCATCTCCTACATCAACATCATGCCTTCGGGTGTTGGCACCATCTGC  
 361 CTCCTGGGATCATCGGGAAACTCCACGGTCATCTTCGGGGTGAAGAAGTCCAAGCTG  
 421 CACTGGTGCAACAAGTCCCCGACATCTCATCATCAACCTCTGGTAGTAGATCTCC  
 481 TTCTCTCTGGCATGCCCTCATGATCCACCAAGCTCATGGCAATAGTCAGTTCACCGACC  
 541 GGGGAGACCATGTCGACCCCTCATCACGGGCCATGGATGCCAATAGTCAGCTAACCT  
 601 TACATCCTGACCGCCATTGACCGCTACCTGGCCACTGGCACTGGCCACCC  
 661 AGGAAGTTCCGGAAAGCCCTCTGGGCCACCCCTGGTATCTGGCTCTGGGCC  
 721 TTCAATCAGCATCACCCCTGTGTGGCTGTATGCCAGACTCATCCCCTCCAGGAGGTGCA  
 781 GTGGGCTGGGATACGGCTGGCCAAACCCAGACACTGACCTCTACTGGTCAACCTGTAC  
 841 CAGTTTTCTGGCCCTTGCCCTGGCTTGTGGTCACTCACAGCCGATAACGTGGGGATC  
 901 CTGCAGGGCATGACGGCTCAGTGGCCCTGGCTGGCTGGCTGGGGACAA  
 961 AAGAGGGTACCCGACAGCCATGCCCATCTGTCTGGTCTGGCTGGGCACCC  
 1021 TACTATGTCAGCTGACCCAGTGGTCCATCAGCCGGCCGACCCCTCACCTTGTCTAC  
 1081 TTATACAATGCCCATCAGCTGGCTATGCCAACAGCTGGCTGGCTGGAGGCCAG  
 1141 ATCGTGCTCTGTGAGACGTTCCGCAAACGCTGGTCTGGTCAAGGGCTGACGAGG  
 1201 GGGCAGGCTTCGGCTGTCAAGAACGCTGACGGGCTGACGAGGAGGAGCAAGAAA  
 1261 GGCACCTGA

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## FIGURE 2

1	M S V G A M K K G V G R A V G L G G G S	20
21	G C Q A T E E D P L P D C G A C A P G Q	40
41	G G R R W R L P Q P A W V E G S S A R L	60
61	W E Q A T G T G W M D L E A S L L P T G	80
81	P N A S N T S D G P D N L T S A G S P P	100
101	R T G S I S Y I N I I M P S V F G T I C	120
121	L L G I I G N S T V I F A V V K K S K L	140
141	H W C N N V P D I F I I N L S V V D L L	160
161	F L L G M P F M I H Q L M G N G V W H F	180
181	G E T M C T L I T A M D A N S Q F T S T	200
201	Y I L T A M A I D R Y L A T V H P I S S	220
221	T K F R K P S V A T L V I C L L W A L S	240
241	F I S I T P V W L Y A R L I P F P G G A	260
261	V G C G I R L P N P D T D L Y W F T L Y	280
281	Q F F L A F A L P F V V I T A A Y V R I	300
301	L Q R M T S S V A P A S Q R S I R L R T	320
321	K R V T R T A I A I C L V F F V C W A P	340
341	Y Y V L Q L T Q L S I S R P T L T F V Y	360
361	L Y N A A I S L G Y A N S C L N P F V Y	380
381	I V L C E T F R K R L V L S V K P A A Q	400
401	G Q L R A V S N A Q T A D E E R T E S K	420
421	G T	422

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## FIGURE 3

1 M S V G A M K K G V G R A V G L G G G S 20  
21 G C Q A T E E D P L P D C G A C A P G Q 40  
41 G G R R W R L P Q P A W V E G S S A R L 60  
61 W E Q A T G T G W M D L E A S L L P T G 80  
81 P N A S N T S D G P D N L T S A G S P P 100  
101 R T G S I S Y I N I IM P S V F G T I C 120  
121 IL L G I I G N S T V I F A V V K K S K L 140  
141 H W C N N V P D IF I I N I S V V D L L 160  
161 E L L G M P F M I H Q L M G N G V W H F 180  
181 G E T M C T L I T A M D A N S O F T S T 200  
182 III  
201 Y I L T A M A I D R Y L A T V H P I S S 220  
221 T K F R K P S V A T L V I C L L W A L S 240  
241 IV  
E I S I T P V W L Y A R L I P F P G G A 260  
261 V G C G I R L P N P D T D L Y W E T L Y 280  
262 V  
281 O F F L A F A L P F V V I T A A Y V R I 300  
301 VI  
301 L Q R M T S S V A P A S Q R S I R L R T 320  
321 K R V T R T A I A I C L V F F V C W A P 340  
341 V Y V L O L T O L S I S R P T L T F V Y 360  
342 VII  
361 L Y N A A I S L G Y A N S C L N P F V Y 380  
381 I V L C E T F R K R L V L S V K P A A Q 400  
401 G Q L R A V S N A Q T A D E E R T E S K 420  
421 G T 422

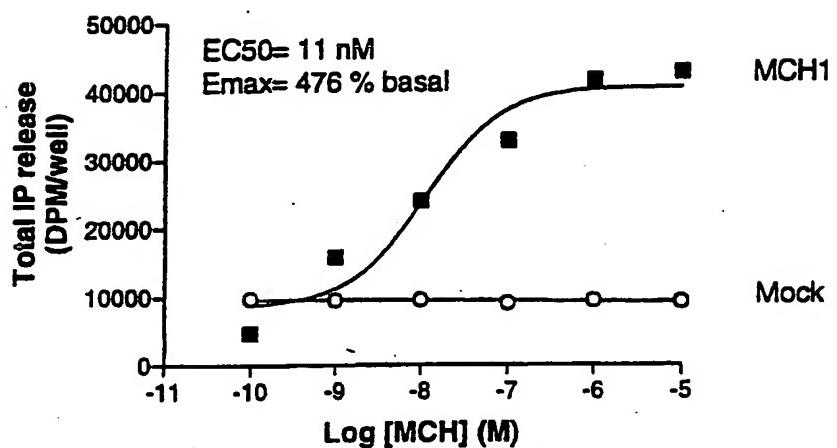
FIGURE 4

1	GCAAGGGACACTGGCACCCGGCTGGCATGGATACTGCAAACCTCGTGTCCACTGGCCCCAA
61	TGCCAGAACATCTCCGATGGCAGGATAATCTCACATTGCCCTGGTCAACCTCCCTCGCAC
121	AGGGAGTGTCTCCATCACATCAACATCATTATGCCCTGGTGGTACCATCTGTCTCCT
181	GGGCATCGTGGAAACTCCACGGTCATCTTGCTGGTGAAGGAAGTCCAAAGCTACACTG
241	GTGCAGCAACGTCGGCACATCTTCATCATCAACCTCTGTGGGATCTGCTCTTCCT
301	GCTGGGCATGCCCTCATGATCACCCAGCTCATGGGGAACCGGGTCTGGCACTTGGGA
361	AACCATGTGCACCCCTCATCACAGGCCATGGACGCCAACAGTCAGTTCACTAGCACCTACAT
421	CCTGACTGCCATGACCATTTGACCGCTACTTGGCCACCGTCCACCCCATCTCCACCAA
481	GTTCGGGAPGCCCTCATGGCCACCGTGGGTGATCTGCCTCCCTGGGGCTCTCCCTCAT
541	CAGTATCACCCCTGTGGCTCATACGCCAGGCTCATTCCTCCAGGGGTGCTGTGGG
601	CTGGGGCATGGCTGGCAAACCGGACACTGACCTCTACTGGTTCACTCTGTACAGTT
661	TTTCCTGGCCTTTCGGCTTGGGTGTCATTACCGCCATACGTGAAATACTACA
721	GCGCATGACGGTCTTCGGTGGCCCAGCCTCCCAACGCAGCATCCGGCTCTGGACAAAGAG
781	GGTACCCGGCACGGCATTGCCATCTGTCTGGTCTCTTGTGTGGCCACCCCTACTA
841	TGGCTGCCAGCTGACCCAGCTGGCCATCAGCCGCCGACCCCTCACGGTTGTCTACTTGT
901	CAACGGGCCATCAGCTGGCTATGCTAACAGCTGGCTGAACCCCTTGTGTACATAGT
961	GCTCTGTGAGACCTTCGAAAACGCTTGGTGTCACTGAAGCCTGGCCAGGGCA
1021	GCTCCGGCACGGTCAGAACGCTCAGACAGCTGATGAGGAGGGACAGAARGCAAGGCAC
1081	CTGACAATCCCCAGTCGGCTCCAAGTCAGGCCACCCATCAAACCGTGGGGAGAGATA
1141	TGAGATTAACCCAAAGGCTTACCCCTGGGAGAATGGAGGGCTGGAGGGCTGGGGCTGTAG
1201	CAACCCACATTCCAC

## FIGURE 5

1	M D L Q T S L L S T G P N A S N I S D G	20
21	Q D N L T L P G S P P R T G S V S Y I N	40
41	I I M P S V F G T I C L L G I V G N S T	60
61	V I F A V V K K S K L H W C S N V P D I	80
81	F I I N L S V V D L L F L L G M P F M I	100
101	H Q L M G N G V W H F G E T M C T L I T	120
121	A M D A N S Q F T S T Y I L T A M T I D	140
141	R Y L A T V H P I S S T K F R K P S M A	160
161	T L V I C L L W A L S F I S I T P V W L	180
181	Y A R L I P F P G G A V G C G I R L P N	200
201	P D T D L Y W F T L Y Q F F L A F A L P	220
221	E V V I T A A Y V K I L Q R M T S S V A	240
241	P A S Q R S I R L R T K R V T R T A I A	260
261	I C L V F F V C W A P Y Y V L Q L T Q L	280
281	S I S R P T L T F V Y L Y N A A I S L G	300
301	Y A N S C L N P F V Y I V L C E T F R K	320
321	R L V L S V K P A A Q G Q L R T V S N A	340
341	Q T A D E E R T E S K G T *	354

FIGURE 6

**IP release in MCH1- and  
mock-transfected Cos-7 cells**

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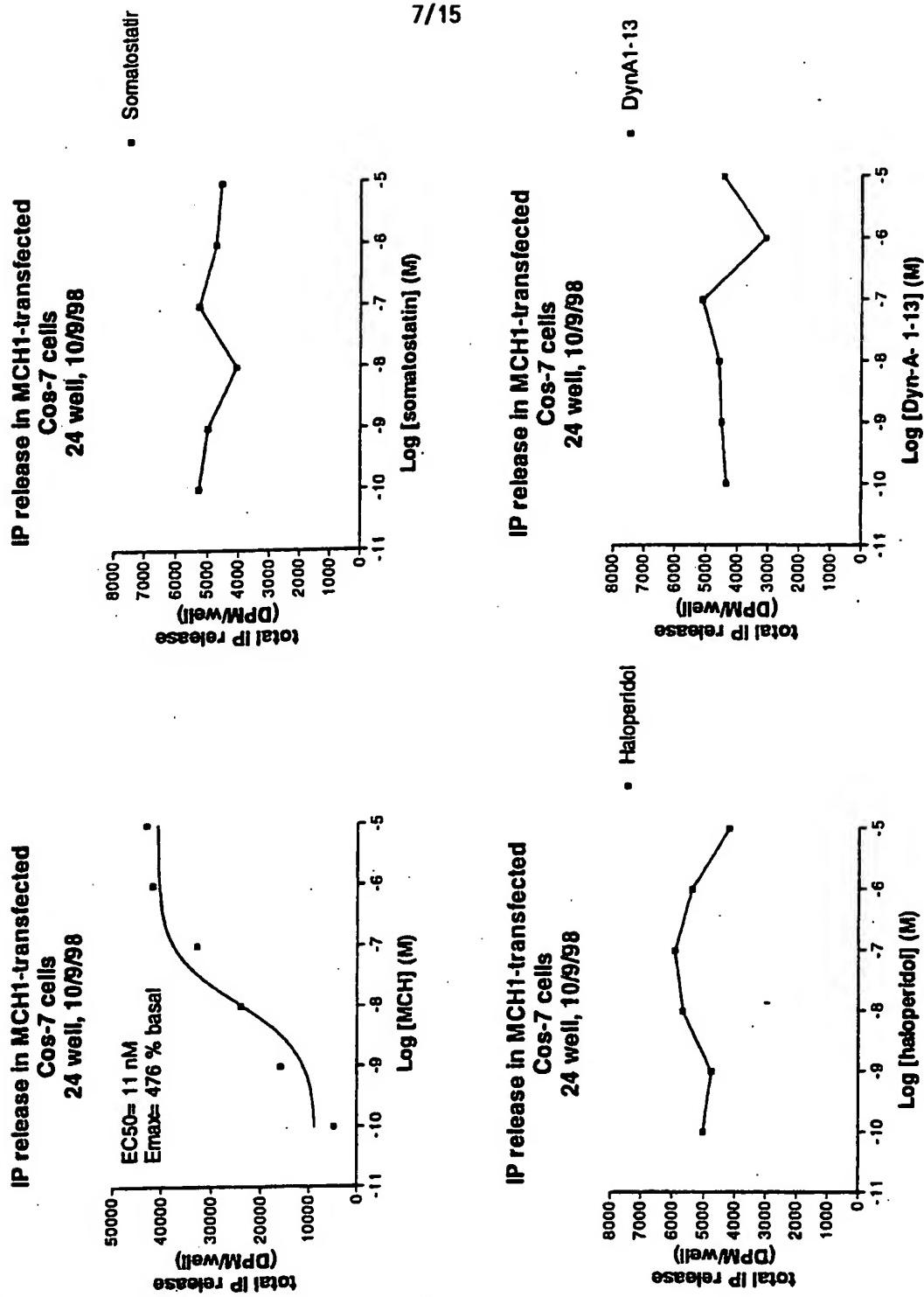


FIGURE 7

**Microphysiometer Response  
CHO cells**

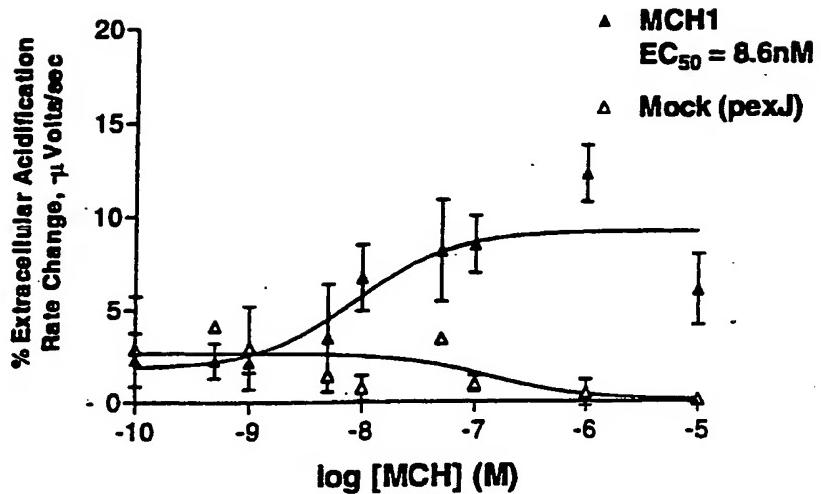
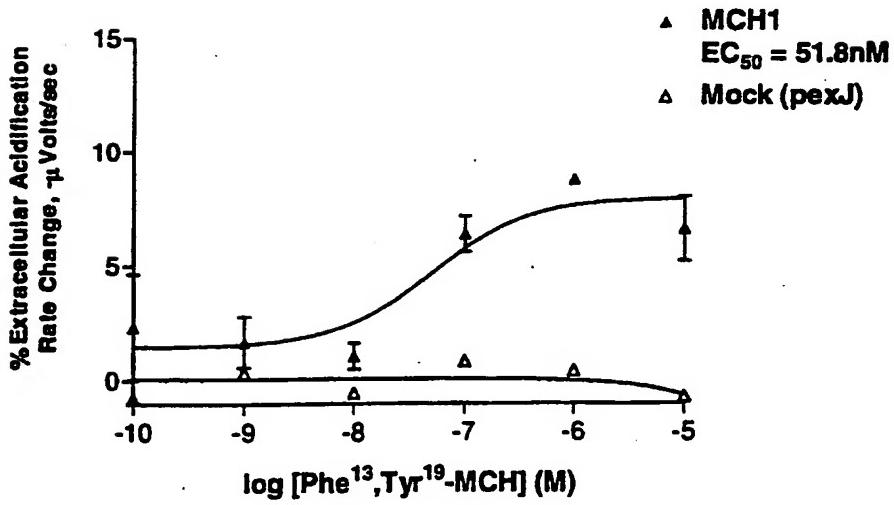


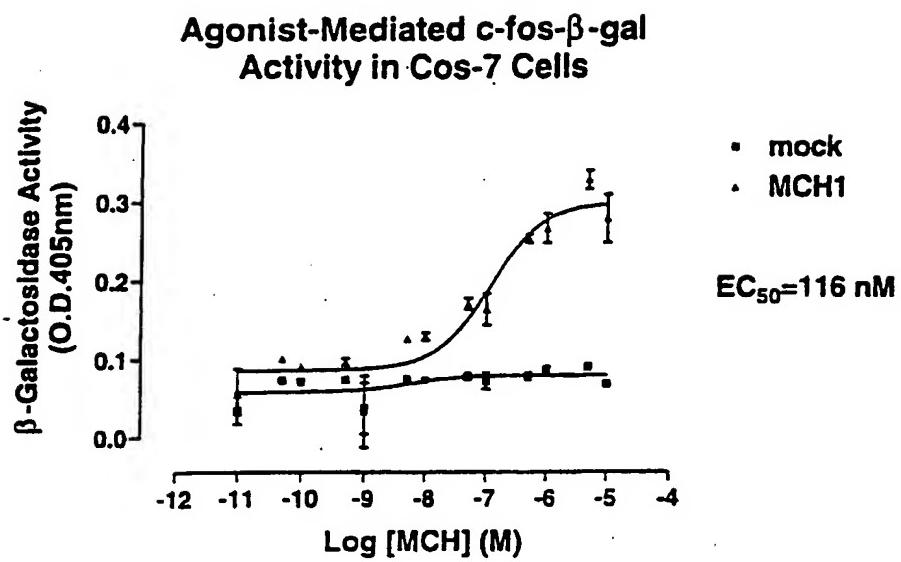
FIGURE 8

**Microphysiometer Response  
CHO cells**



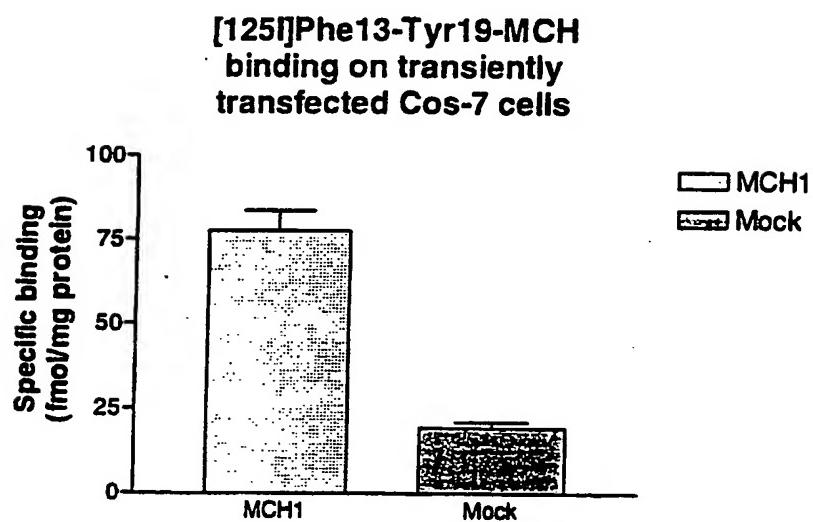
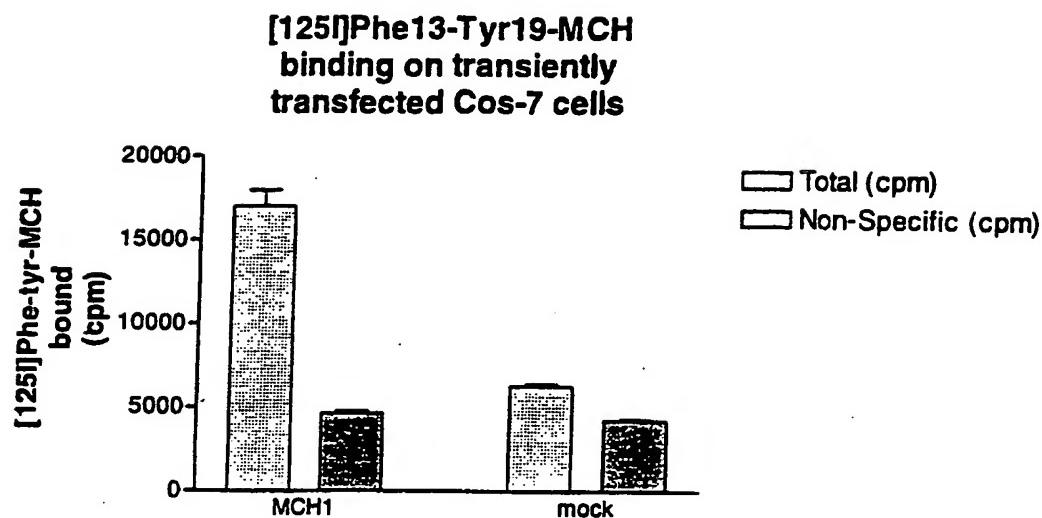
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FIGURE 9



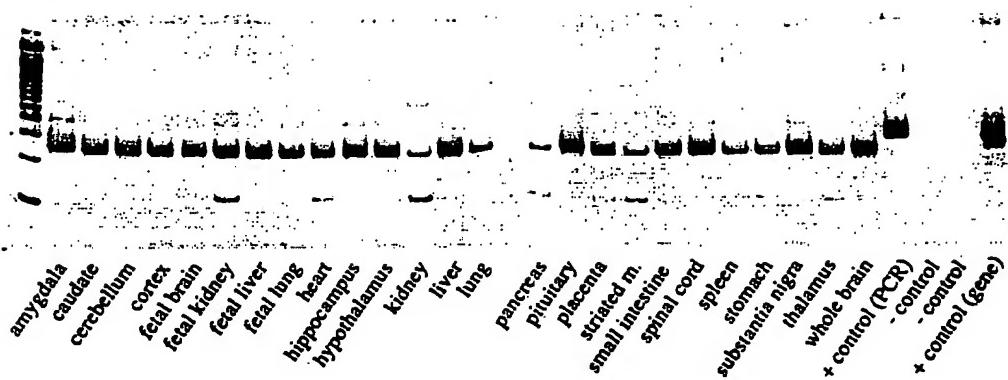
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FIGURE 10



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## FIGURE 11



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Figure 12

	1		40
TL231	MSVGAMKKGV	GRAVGLGGGS	GCQATEEDPL PDCGACAPGQ
R106	MSVGAMKKGV	GRAVGLGGGS	GCQATEEDPL PDCGACAPGQ
R114	MSVGAaKKGV	GRAVGLGGGS	GCQATEEDPL PDCGACAPGQ
BO120	~~~~~	~~~~~	~~~~~

	41		80
TL231	GGRRWRLPQP	AWVEGSSARL	WEQATGTGWM DLEASLLPTG
R106	GGRRWRLPQP	AWVEGSSARL	WEQATGTGWA DLEASLLPTG
R114	GGRRWRLPQP	AWVEGSSARL	WEQATGTGWA DLEASLLPTG
BO120	~~~~~	~~~~~	~~~~~M DLEASLLPTG

	81		100
TL231	PNASNTSDGP	DNLTSAGSPP...	
R106	PNASNTSDGP	DNLTSAGSPP...	
R114	PNASNTSDGP	DNLTSAGSPP...	
BO120	PNASNTSDGP	DNLTSAGSPP...	

1	M S V G A M K K G V G R A V G L G G G S	20
21	G C Q A T E E D P L P D C G A C A P G Q	40
41	G G R R W R L P Q P A W V E G S S A R L	60
61	W E Q A T G T G W A D L E A S L L P T G	80
81	P N A S N T S D G P D N L T S A G S P P	100
101	R T G S I S Y I N I I M P S V F G T I C	120
121	L L G I I G N S T V I F A V V K K S K L	140
141	H W C N N V P D I F I I N L S V V D L L	160
161	F L L G M P F M I H Q L M G N G V W H F	180
181	G E T M C T L I T A M D A N S Q F T S T	200
201	Y I L T A M A I D R Y L A T V H P I S S	220
221	T K F R K P S V A T L V I C L L W A L S	240
241	F I S I T P V W L Y A R L I P F P G G A	260
261	V G C G I R L P N P D T D L Y W F T L Y	280
281	Q F F L A F A L P F V V I T A A Y V R I	300
301	L Q R M T S S V A P A S Q R S I R L R T	320
321	K R V T R T A I A I C L V F F V C W A P	340
341	Y Y V L Q L T Q L S I S R P T L T F V Y	360
361	L Y N A A I S L G Y A N S C L N P F V Y	380
381	I V L C E T F R K R L V L S V K P A A Q	400
401	G Q L R A V S N A Q T A D E E R T E S K	420
421	G T	422

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1	M S V G A A K K G V G R A V G L G G G S	20
21	G C Q A T E E D P L P D C G A C A P G Q	40
41	G G R R W R L P Q P A W V E G S S A R L	60
61	W E Q A T G T G W A D L E A S L L P T G	80
81	P N A S N T S D G P D N L T S A G S P P	100
101	R T G S I S Y I N I I M P S V F G T I C	120
121	L L G I I G N S T V I F A V V K K S K L	140
141	H W C N N V P D I F I I N L S V V D L L	160
161	F L L G M P F M I H Q L M G N G V W H F	180
181	G E T M C T L I T A M D A N S Q F T S T	200
201	Y I L T A M A I D R Y L A T V H P I S S	220
221	T K F R K P S V A T L V I C L L W A L S	240
241	F I S I T P V W L Y A R L I P F P G G A	260
261	V G C G I R L P N P D T D L Y W F T L Y	280
281	Q F F L A F A L P F V V I T A A Y V R I	300
301	L Q R M T S S V A P A S Q R S I R L R T	320
321	K R V T R T A I A I C L V F F V C W A P	340
341	Y Y V L Q L T Q L S I S R P T L T F V Y	360
361	L Y N A A I S L G Y A N S C L N P F V Y	380
381	I V L C E T F R K R L V L S V K P A A Q	400
401	G Q L R A V S N A Q T A D E E R T E S K	420
421	G T	422

1	M D L E A S L L P T G P N A S N T S D G	20
21	P D N L T S A G S P P R T G S I S Y I N	40
41	I I M P S V F G T I C L L G I I G N S T	60
61	V I F A V V K K S K L H W C N N V P D I	80
81	F I I N L S V V D L L F L L G M P F M I	100
101	H Q L M G N G V W H F G E T M C T L I T	120
121	A M D A N S Q F T S T Y I L T A M A I D	140
141	R Y L A T V H P I S S T K F R K P S V A	160
161	T L V I C L L W A L S F I S I T P V W L	180
181	Y A R L I P F P G G A V G C G I R L P N	200
201	P D T D L Y W F T L Y Q F F L A F A L P	220
221	F V V I T A A Y V R I L Q R M T S S V A	240
241	P A S Q R S I R L R T K R V T R T A I A	260
261	I C L V F F V C W A P Y Y V L Q L T Q L	280
281	S I S R P T L T F V Y L Y N A A I S L G	300
301	Y A N S C L N P F V Y I V L C E T F R K	320
321	R L V L S V K P A A Q G Q L R A V S N A	340
341	Q T A D E E R T E S K G T	353



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 : <b>C07K 14/72, C07H 21/04, C12N 15/12, 15/63, 1/21, C12P 21/02, G01N 33/53</b>		A3	(11) International Publication Number: <b>WO 00/39279</b> (43) International Publication Date: <b>6 July 2000 (06.07.00)</b>
(21) International Application Number: <b>PCT/US99/31169</b> (22) International Filing Date: <b>30 December 1999 (30.12.99)</b>		(74) Agent: <b>WHITE, John, P.; Cooper &amp; Dunham LLP, 1185 Avenue of the Americas, New York, NY 10036 (US).</b>	
(30) Priority Data: <b>09/224,426 31 December 1998 (31.12.98) US</b>		(81) Designated States: <b>AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</b>	
(63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application <b>US 09/224,426 (CIP) Filed on 31 December 1998 (31.12.98)</b>		Published <i>With international search report.</i>	
(71) Applicant (for all designated States except US): <b>SYNAPTIC PHARMACEUTICAL CORPORATION [US/US]; 215 College Road, Paramus, NJ 07652 (US).</b>		(88) Date of publication of the international search report: <b>2 November 2000 (02.11.00)</b>	
(54) Title: <b>DNA ENCODING A HUMAN MELANIN CONCENTRATING HORMONE RECEPTOR (MCH1) AND USES THEREOF</b>			
(57) Abstract			
<p>This invention provides an isolated nucleic acid encoding a human MCH1 receptor, a purified human MCH1 receptor, vectors comprising isolated nucleic acid encoding a human MCH1 receptor, cells comprising such vectors, antibodies directed to a human MCH1 receptor, nucleic acid probes useful for detecting nucleic acid encoding human MCH1 receptors, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding human MCH1 receptors, transgenic, nonhuman animals which express DNA encoding a normal or mutant human MCH1 receptor, methods of isolating a human MCH1 receptor, methods of treating an abnormality that is linked to the activity of a human MCH1 receptor, as well as methods of determining binding of compounds to mammalian MCH1 receptors.</p>			

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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US99/31169

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(7) :Please See Extra Sheet.

US CL :Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. :

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 848 060 A2 (SMITHKLINE BEECHAM CORPORATION) 17 June 1998 (17.06.98), see entire document, especially SEQ ID NOS: 1 and 2, claims 1-11 and 20, and pages 3, 4, 8-10 and 14-16.	1-5, 10-15, 17-23, 25-28, 47, 49, 51, 52, 56, 57, 59-63, 65, 67, 69-73, 97-100, 106-110 and 112-116  6-9, 16, 48, 50, 53-55, 66, 68, 75-81, 101, 111, 122-130 and 147-154
A		

 Further documents are listed in the continuation of Box C.  See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E* earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"a."	document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means		
*P* document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

06 JUNE 2000

Date of mailing of the international search report

13 JUL 2000

Name and mailing address of the ISA/US  
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EILEEN O'HARA

Telephone No. (703) 308-0196

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US99/31169

**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-23,25-28,47-57,59-63,65-73,75-81,97-101,106-116,122-130,147-154

Remark on Protest



The additional search fees were accompanied by the applicant's protest.



No protest accompanied the payment of additional search fees.

**INTERNATIONAL SEARCH REPORT**

International application No. PCT/US99/31169
---

**A. CLASSIFICATION OF SUBJECT MATTER:  
IPC (7):**

C07K 14/72; C07H 21/04; C12N 15/12, 15/63, 1/21; C12P 21/02; G01N 33/53

**A. CLASSIFICATION OF SUBJECT MATTER:  
US CL :**

530/350; 536/23.5, 24.31; 435/6, 7.1, 7.2, 69.1, 252.3, 254.2, 361

**BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING**  
This ISA found multiple inventions as follows:

This application contains the following inventions or groups of inventions which are not so linked as to form a single inventive concept under PCT Rule 13.1. In order for all inventions to be searched, the appropriate additional search fees must be paid.

Group I, claim(s) 1-23, 25-28, 47-57, 59-63, 65-73, 75-81, 97-101, 106-116, 122-130, 147-154, drawn to nucleic acids encoding MCH1 receptors, vectors, host cells, method of recombinantly producing MCH1 receptor protein, and methods for identifying compounds that bind to or activate or inhibit MCH1 receptor.

Group II, claim(s) 24, drawn to a membrane preparation.

Group III, claims 29-31 and 35-39, drawn to antisense oligonucleotides.

Group IV, claims 32, 34 and 40, drawn to antibodies.

Group V, claim 33, drawn to an agent capable of competitively inhibiting binding of antibody to MCH1 receptor.

Group VI, claims 41, 44, 45 and 46, drawn to transgenic mammals comprising human MCH1 receptor.

Group VII, claims 42, 44, 45 and 46, drawn to transgenic mammals having an MCH1 knockout.

Group VIII, claims 43 and 46, drawn to transgenic mammals comprising antisense DNA complementary to MCH1 DNA.

Group IX, claim 82, drawn to a method of detecting expression of MCH1 receptor by northern hybridization.

Group X, claim 83, drawn to a method of detecting MCH1 receptor protein on the surface of a cell by antibody binding.

Group XI, claims 84 and 85, drawn to a method of determining the physiological effects of varying levels of MCH1 receptor in a transgenic mammal comprising an inducible promoter regulating MCH1 expression.

Group XII, claims 86 and 90, drawn to a method of indentifying antagonists or agonists of MCH1 receptor in a transgenic animal expressing the human MCH1 receptor.

Group XIII, claims 58, 64, 74, 87, 88, 104, 105, 117, 120, 121 and 132, drawn to antagonists of MCH1 receptor.

Group XIV, claims 89, 135 and 136, drawn to a method of treatment by administering an antagonist to MCH1.

Group XV, claims 58, 64, 74, 91, 92, 102, 103, 117, 118, 119 and 131, drawn to agonists of MCH1 receptor.

Group XVI, claims 93, 133 and 134, drawn to a method of treatment by administration of an agonist to MCH1 receptor.

Group XVII, claims 94 and 95, drawn to a method of diagnosis of a disorder by restriction fragment length polymorphism.

Group XVIII, claim 96, drawn to a method of preparing purified MCH1 receptor.

Group XIX, claims 137-146 and 155-168, drawn to a process for making a composition of matter.

The inventions listed as Groups I-XIX do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Pursuant to 37 C.F.R. § 1.475(d), the ISA/US considers that where multiple products and processes are claimed, the main invention shall consist of the first invention of the category first mentioned in the claims and the first recited invention of each of the other categories related thereto. Accordingly, the main invention (Group I) comprises the first recited product, polynucleotides encoding MCH1 receptor, host cells, vectors, a recombinant method of producing MCH1 receptor protein, the MCH1 protein, and methods for identifying compounds that bind to or modulate MCH1 receptor activity. Further pursuant to 37 C.F.R. § 1.475(d), the ISA/US considers that any feature which the subsequently recited products and methods share with the main invention does not constitute a special technical feature within the meaning of PCT Rule 13.2, and that each of such products and methods accordingly defines a separate invention.